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(71) Applicant: SCHERING CORPORATION [US/US]; 2000 Galloping Hill Road, Kenilworth, NJ 07033 (US).

(72) Inventors: SHUE, Ho-Jane; 8 Hamilton Place, Pine Brook, NJ 07058 (US). SHIH, Neng-Yang; 1 Maple Drive, North Caldwell, NJ 07006 (US). BLYTHIN, David, J.; 487 Mountain Avenue, North Caldwell, NJ 07006 (US). CHEN, Xiao; 15 Hillwood Avenue, Edison, NJ 08820 (US). PIWINSKI, John, J.; 6 Saddle Ridge Drive, Clinton Township, NJ 08833 (US). MCCORMICK, Kevin, D.; 5 Pace Drive, Edison, NJ 08820 (US).

(74) Agents: MAGATTI, Anita, W. et al.; Schering-Plough Corporation, Patent Dept. K-6-1 1990, 2000 Galloping Hill Road, Kenilworth, NJ 07033 (US).

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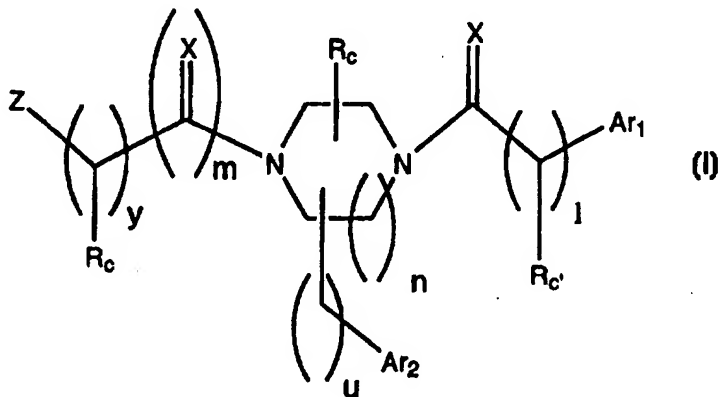
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(54) Title: PIPERAZINO DERIVATIVES AS NEUROKININ ANTAGONISTS

(57) Abstract

The invention relates to compounds of formula (I). These compounds are neurokinin antagonists. These compounds are useful in the treatment of chronic airway diseases such as asthma.



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5 PIPERAZINO DERIVATIVES AS NEUROKININ ANTAGONISTS

BACKGROUND OF THE INVENTION

The present invention relates to a genus of compounds useful as antagonists of neurokinin receptors. In particular, these can be
10 neurokinin-1 receptor (NK₁) antagonists. Some can also be neurokinin-1 receptor (NK₁) antagonists and neurokinin-2 receptor (NK₂) antagonists, that is, NK₁ / NK₂ dual receptor antagonists. Some can also be neurokinin-2 receptor (NK₂) antagonists. Some can also be neurokinin-3 receptor (NK₃) antagonists.

15 Neurokinin receptors are found in the nervous system and the circulatory system and peripheral tissues of mammals, and therefore are involved in a variety of biological processes. Neurokinin receptor antagonists are consequently expected to be useful in the treatment or prevention of various mammalian disease states, for example pulmonary
20 disorders like asthma, cough, bronchospasm, chronic obstructive pulmonary diseases, and airway hyperreactivity; skin disorders and itch, for example, atopic dermatitis, and cutaneous wheal and flare; neurogenic inflammation inflammatory diseases such as arthritis, migraine, nociception; CNS diseases such as anxiety, Parkinson's disease,
25 movement disorders and psychosis; convulsive disorders, renal disorders, urinary incontinence, ocular inflammation, inflammatory pain, and eating disorders such as food intake inhibition; allergic rhinitis, neurodegenerative disorders, psoriasis, Huntington's disease, depression, emesis and various gastrointestinal disorders such as
30 Crohn's disease.

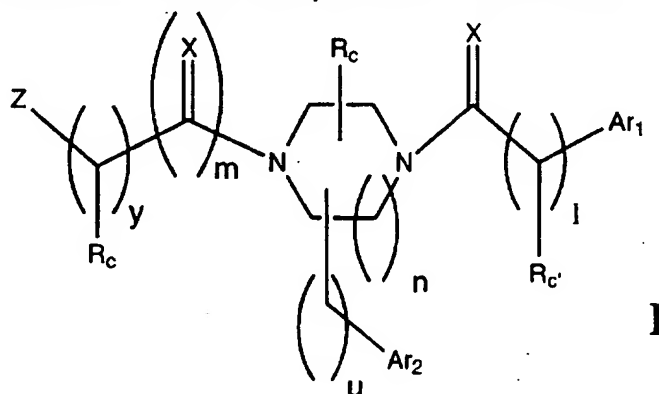
In particular, NK₁ receptors have been reported to be involved in microvascular leakage and mucus secretion, and NK₂ receptors have been associated with smooth muscle contraction, making NK₁ and NK₂ receptor antagonists especially useful in the treatment and
35 prevention of asthma.

Moreover, NK₃ receptor antagonists are especially useful in the treatment and prevention of asthma, inflammatory diseases and conditions, such as ocular inflammation, allergic rhinitis, cutaneous wheal

and flare, psoriasis, atopic dermatitis, CNS diseases such as anxiety and Parkinson's disease.

Summary of the Invention

5 The invention relates to compounds of the formula:

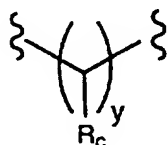


each X is independently, O , (H,H) , NR_d , or S ;

n is 0 to 2; u is 0 to 2; l is 0 to 2;

m is 1, and y is 1 to 3; or m is 2, and y is 0;

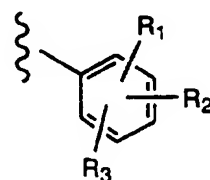
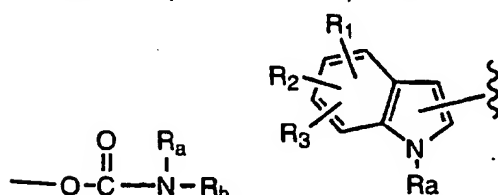
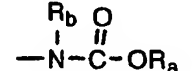
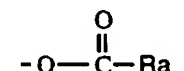
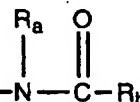
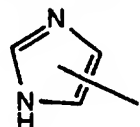
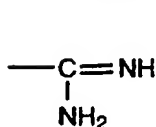
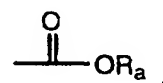
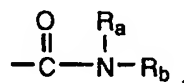
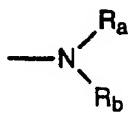
10 each R_C is independently H, C₁-C₆ alkyl, -(CH₂)_{n1}-R₄ where n₁ is 1 to 6;
R_d is independently selected from the group consisting of H, C₁-C₆ alkyl,
CN, OR_a, phenyl, substituted phenyl, benzyl, substituted benzyl, or allyl,
and with the further proviso that no more than one R_C is other than H in the



moiety;

15

R_4 is $-OR_a$, SR_a , $-CN$,

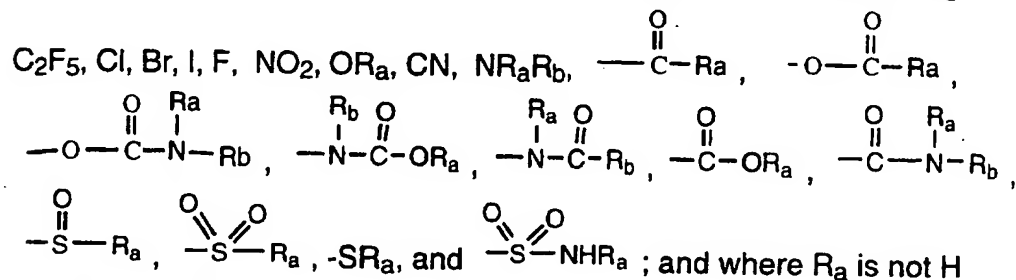


R_c' is H, C_1 - C_6 alkyl or $(CH_2)_nOR_a$, with the proviso that no more than one R_c' is other than H;

- 3 -

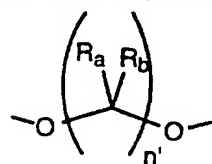
- each R_a and R_b is independently selected from the group consisting of H, C₁-C₆ alkyl, phenyl, substituted phenyl, benzyl, substituted benzyl, allyl; or when R_a and R_b are attached to the same nitrogen, then R_a and R_b together with the nitrogen to which they are attached, form a 4 to 7 member ring;

wherein each R_1 and R_2 is independently H, C₁-C₆ alkyl, CF₃,



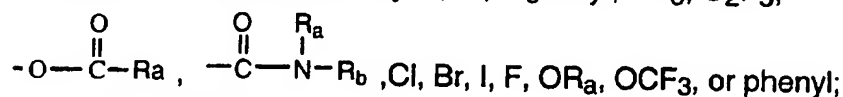
- 10 in $\text{---}\overset{\text{O}}{\parallel}\text{S---Ra}$, $\text{---}\overset{\text{O}}{\parallel}\text{S---}\overset{\text{O}}{\parallel}\text{S---Ra}$, or $\text{---N(Rb)---}\overset{\text{O}}{\parallel}\text{C---ORa}$;

or when R_1 and R_2 are on adjacent carbons on a ring, they can form

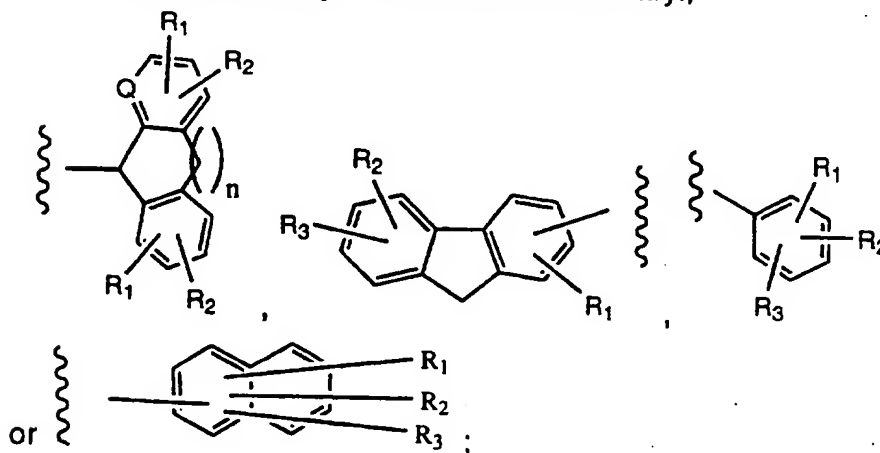


wherein n' is 1 or 2;

and each R_3 is independently H, C₁-C₆ alkyl, CF₃, C₂F₅, $\text{---}\overset{\text{O}}{\parallel}\text{C---Ra}$,



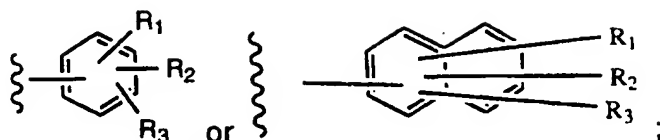
- 15 Ar₁ is heteroaryl or substituted heteroaryl,



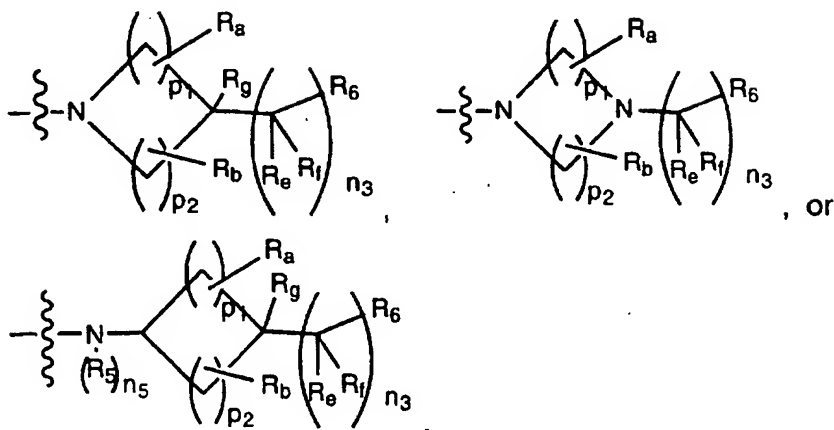
Q is N or CH;

Ar₂ is heteroaryl, substituted heteroaryl,

- 4 -



Z is



- 5 wherein p_1 and p_2 are each independently 1 to 4 with the proviso that p_1 and p_2 added together are 2 to 6; n_5 is 1 to 2;

each R_5 is independently selected from the group consisting of H,

- 10 OH, $\text{—}\overset{\text{O}}{\parallel}\text{C—R}_a$, $\text{C}_1\text{—C}_6$ alkyl, $\text{—}(\text{CH}_2)_{n_1}\text{—R}_4$ where n_1 is 1 to 6 with the proviso that when n_1 is 1, R_4 is not OH or NR_aR_b ; also with the proviso that when n_5 is 2, R_5 is $\text{C}_1\text{—C}_6$ alkyl, and two R_5 can be attached to the nitrogen to form a quaternary salt;

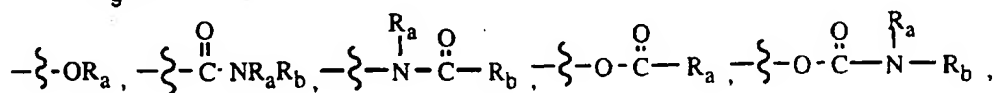
each R_a and R_b is independently selected from the group consisting of H, $\text{C}_1\text{—C}_6$ alkyl, phenyl, substituted phenyl, benzyl, substituted benzyl, allyl;

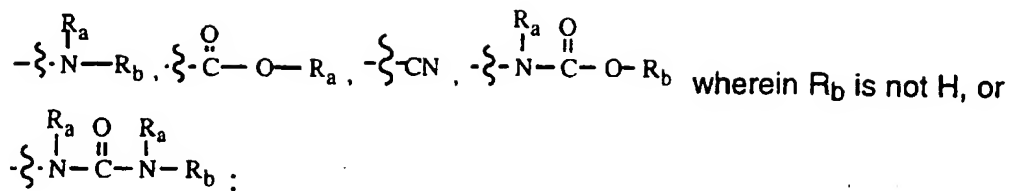
- 15 n_3 is 0-4;

each R_e and R_f is independently selected from the group consisting of H, $\text{C}_1\text{—C}_6$ alkyl, phenyl, substituted phenyl, benzyl, substituted benzyl, allyl; or R_e and R_f taken together with the carbon to which they are attached can also form a carbonyl group with the proviso that no more

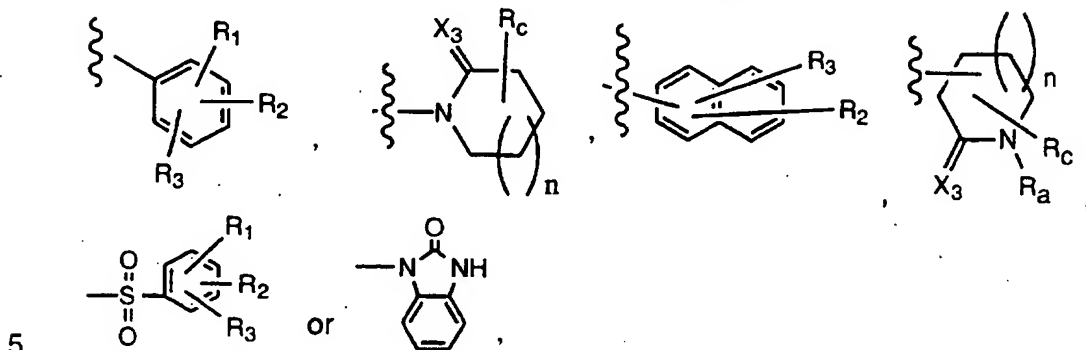
- 20 than one carbonyl group is in the $\text{—}\left(\text{—}\overset{\text{R}_e}{\text{C}}\text{—}\overset{\text{R}_f}{\text{C}}\text{—}\right)_{n_3}\text{—}$ moiety;

R_g is H,

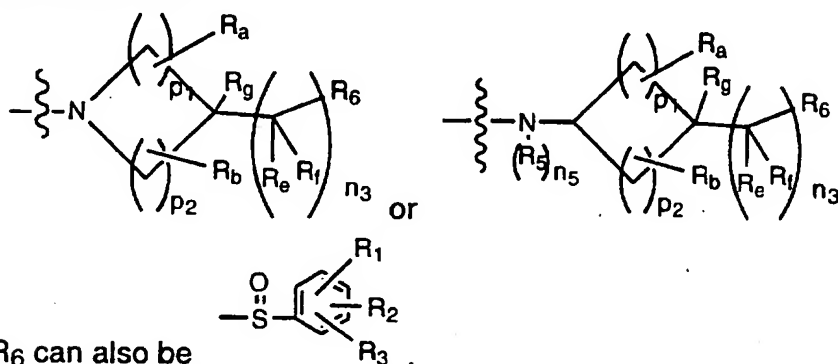




R₆ is H, C₁-C₆ alkyl, allyl, C₃-C₆ cycloalkyl,



and when Z is



R₆ can also be

wherein X_3 is O, (H,H), NR_d , or S; or R_6 is heteroaryl, substituted

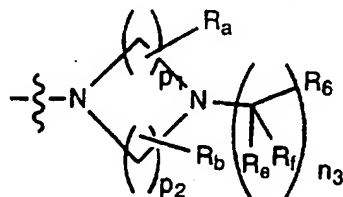
10 heteroaryl, heterocycloalkyl, substituted heterocycloalkyl, when n_3 is 0-4; or when R_e , R_f taken together with the carbon atom to which they are attached form a carbonyl group and n_3 is 1, R_6 can also be OR_a wherein R_a is not H, and R_6 can also be $-(NR_a, R_b)$, O-heteroaryl, O-substituted heteroaryl, O-heterocycloalkyl, O-substituted heterocycloalkyl, $-NR_a$ -
15 heteroaryl, $-NR_a$ -substituted heteroaryl, $-NR_a$ -heterocycloalkyl, $-NR_a$ -substituted heterocycloalkyl.

or a pharmaceutically acceptable salt thereof.

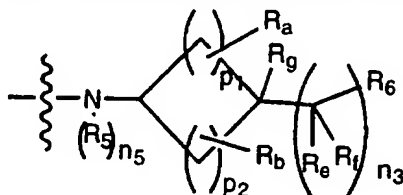
20 All of the variables in the above formulas such as Z, R₁, R₂, and R₃,
have the same meaning throughout the specification unless otherwise
specified.

Preferred compounds of the invention are compounds of formula I, wherein each X is O or (H,H) and at least one X is O.

Also preferred are compounds of formula I wherein Z is

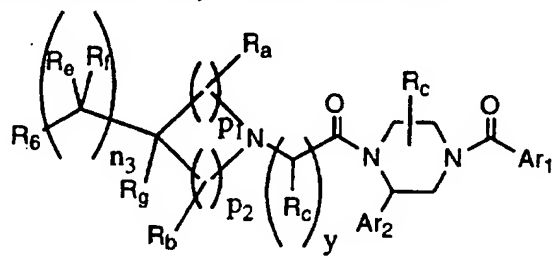


Also preferred are compounds of formula I wherein Z is



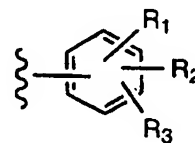
- 5 In the rings above, one R_a and one R_b can be present at any position on the ring which will allow for a substitution.

Also preferred are compounds of formula II



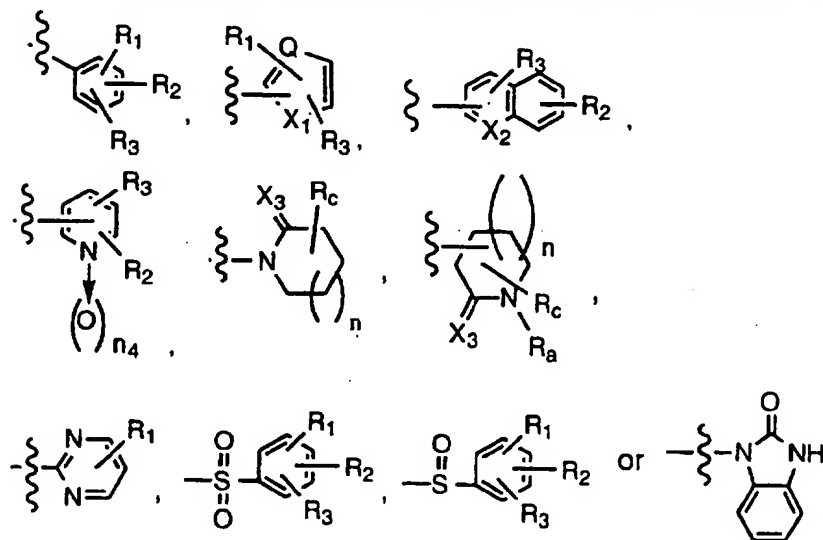
II

wherein R_C is H; y is 1-3; p₁ and p₂ are 2; R_e and R_f are H, C₁-C₆ alkyl, C₃-



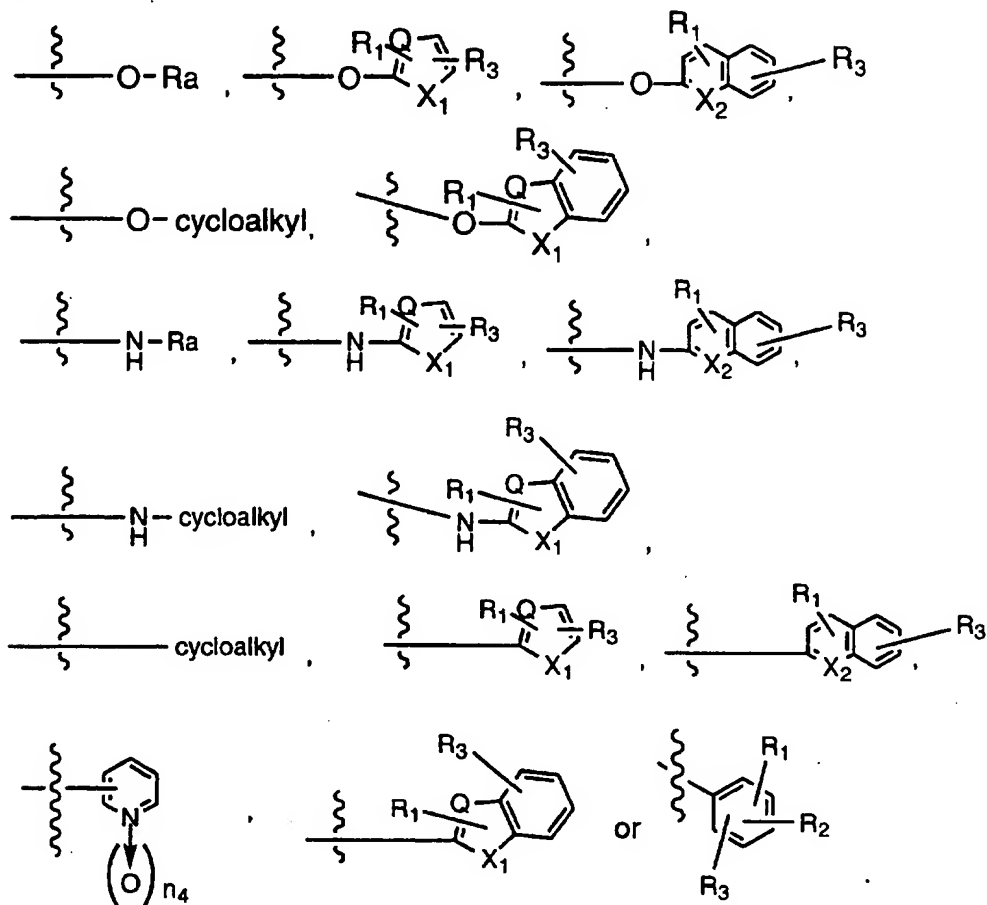
- 10 C₆ cycloalkyl or allyl; n₃ is 0-4; and Ar₁ and Ar₂ are both

Also preferred are compounds of formula II wherein R₆ is

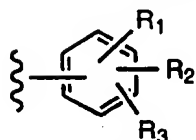


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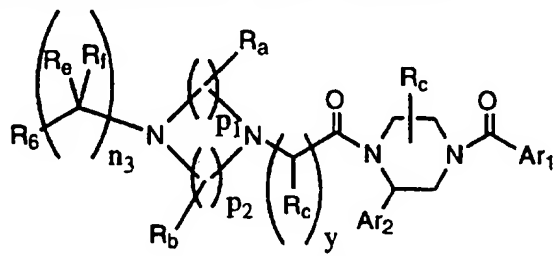
or R_e , R_f taken together with the carbon to which they are attached form a carbonyl group, n_3 is 1 and R_6 is



5 Also preferred are compounds of formula II wherein R_6 is

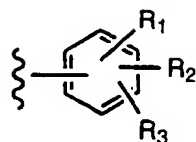


Also preferred are compounds of formula III

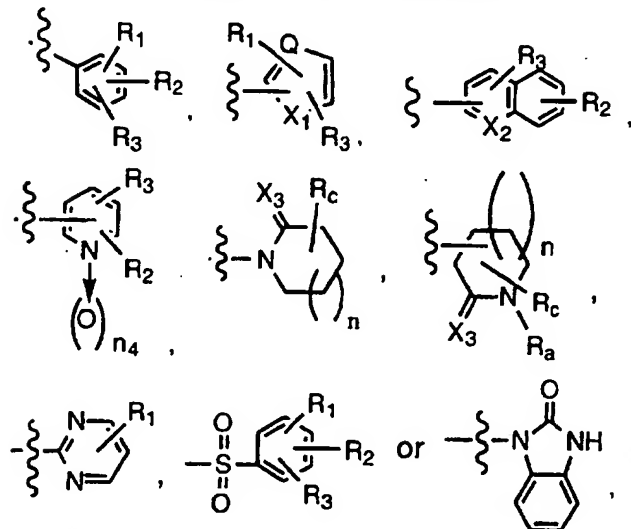


wherein R_c is H; y is 1-3; p_1 and p_2 are 2; R_e and R_f are H, C_1 - C_6 alkyl,

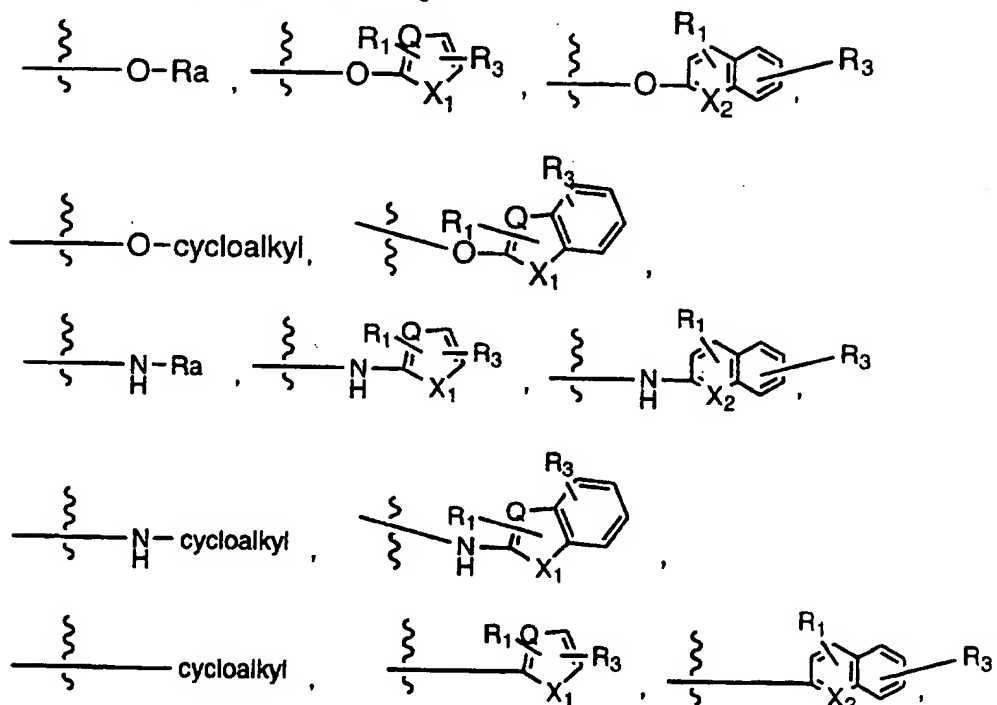
10 C_3 - C_6 cycloalkyl or allyl; n_3 is 0-4; and Ar_1 and Ar_2 are both



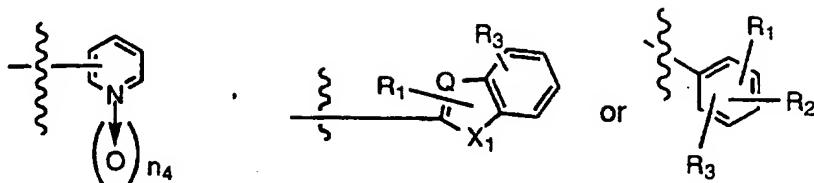
Also preferred are compounds of formula III wherein R_6 is



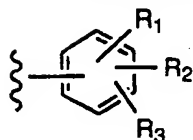
- 5 or R_e , R_f taken together with the carbon to which they are attached form a carbonyl group, n_3 is 1, and R_6 is



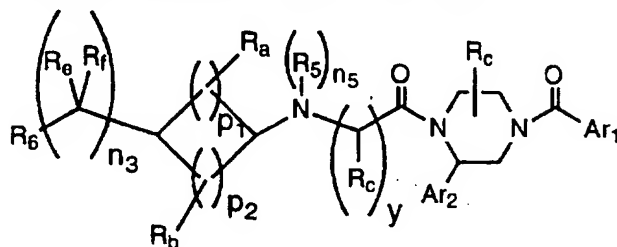
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Also preferred are compounds of formula III wherein R_6 is



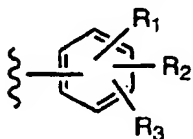
Also preferred are compounds of formula IV



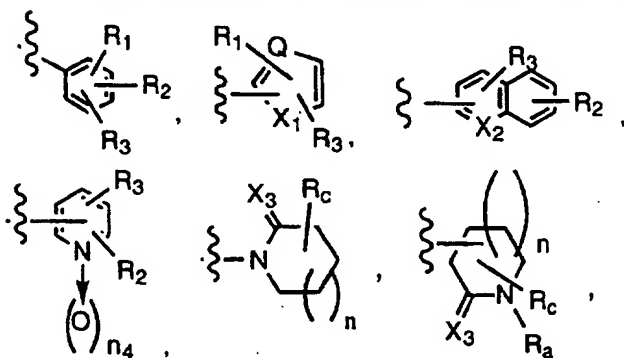
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IV

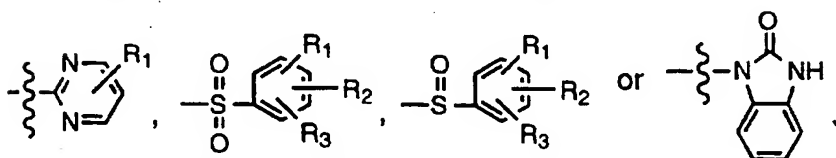
wherein R_c is H; y is 1-3; p_1 , p_2 are 1-2; R_e and R_f are H, C₁-C₆ alkyl, C₃-C₆ cycloalkyl or allyl; n_3 is 0-4; and Ar_1 and Ar_2 are both



Also preferred are compounds of formula IV wherein R_6 is

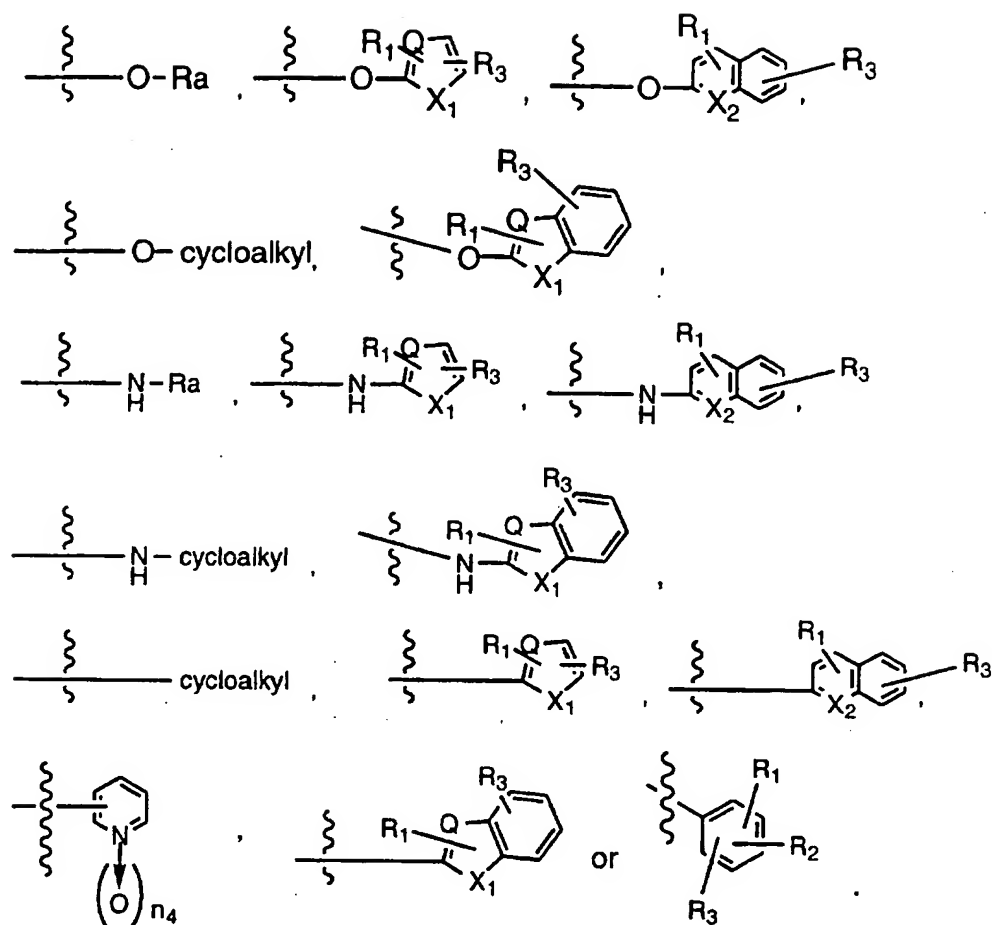


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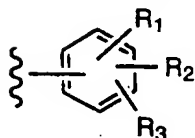


or R_e , R_f taken together with the carbon to which they are attached form a carbonyl group, n_3 is 1, and R_6 is

- 11 -

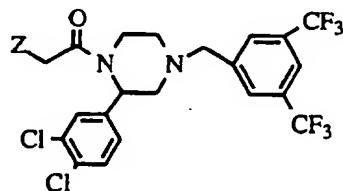


Also preferred are compounds of formula IV wherein R₆ is

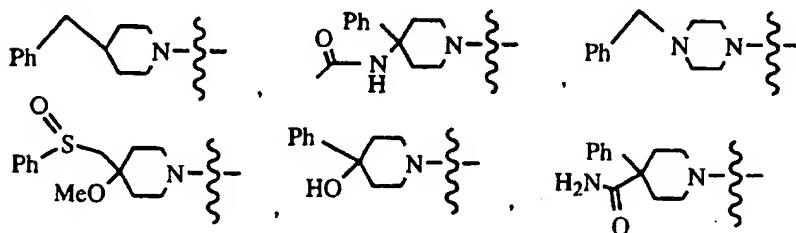


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Exemplary compounds of the invention are compounds of the formulas:

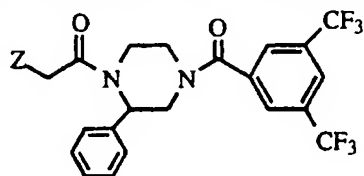


Where Z is

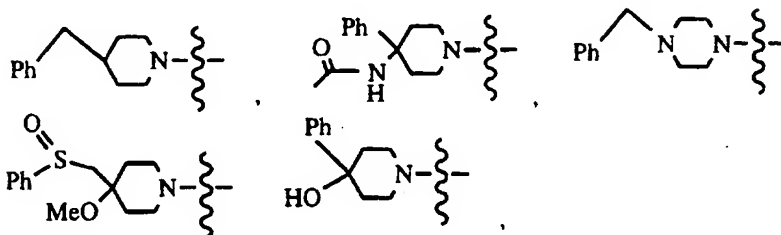


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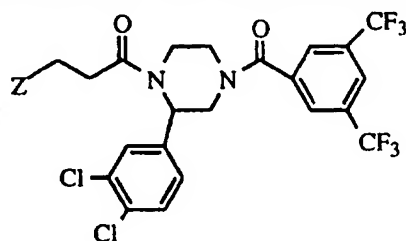
or a compound selected from the group consisting of



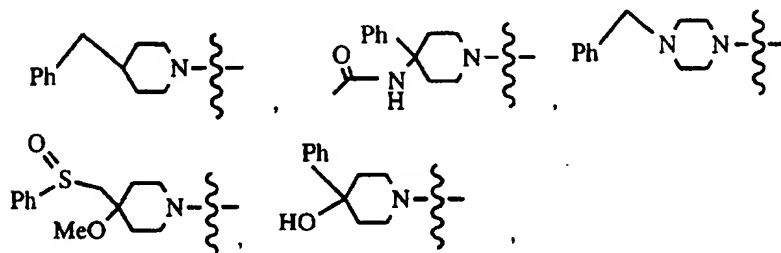
Where Z is



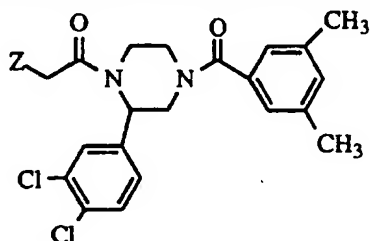
or a compound selected from the group consisting of



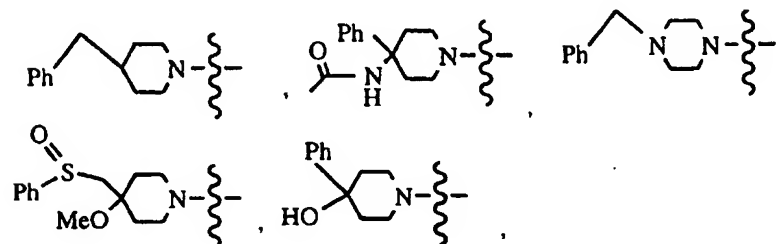
Where Z is



5 or a compound selected from the group consisting of

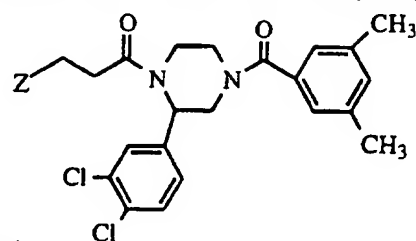


Where Z is

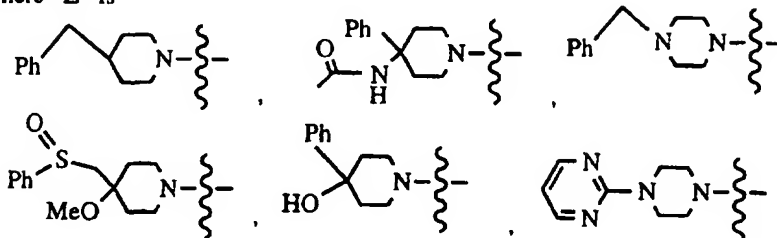


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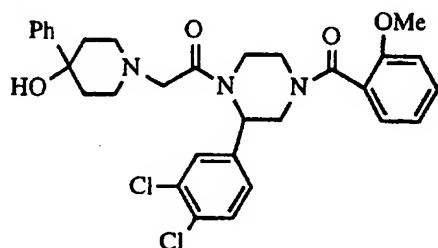
or a compound selected from the group consisting of



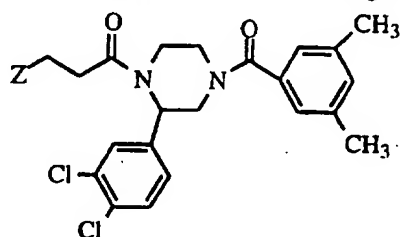
Where Z is



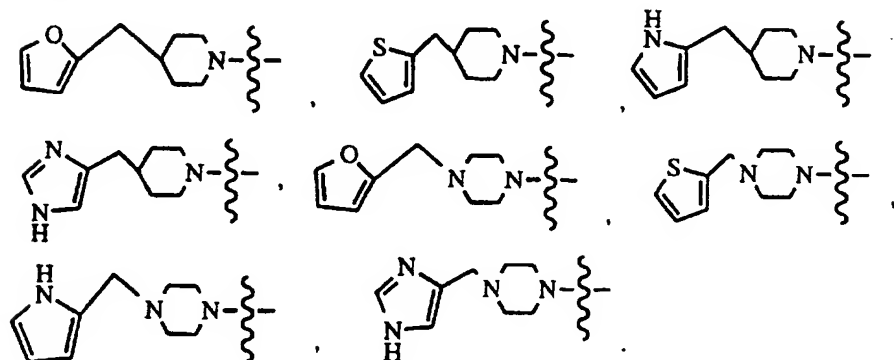
or



5 or a compound selected from the group consisting of

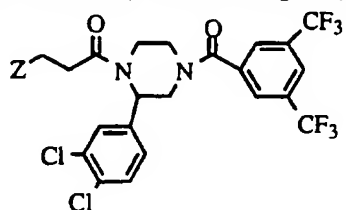


Where Z is

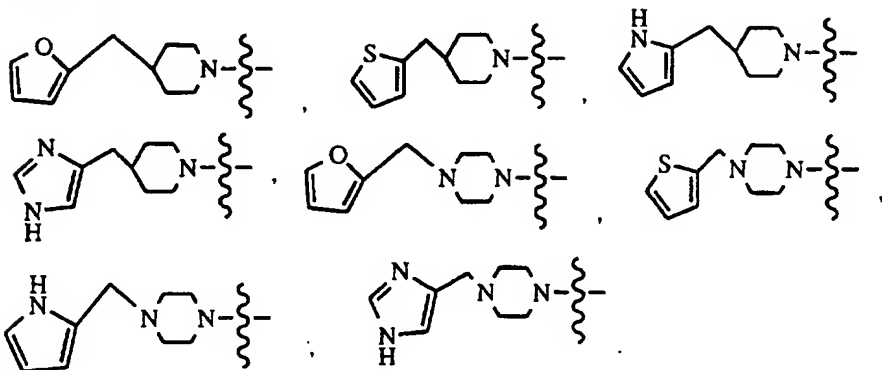


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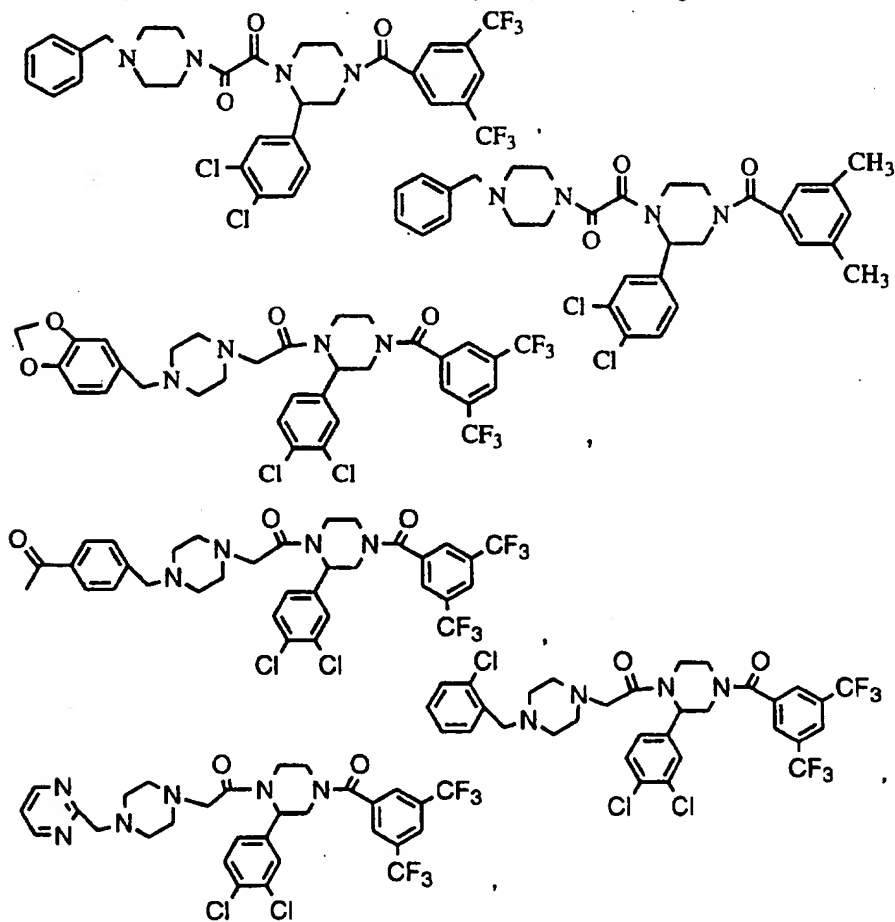
or a compound selected from the group consisting of

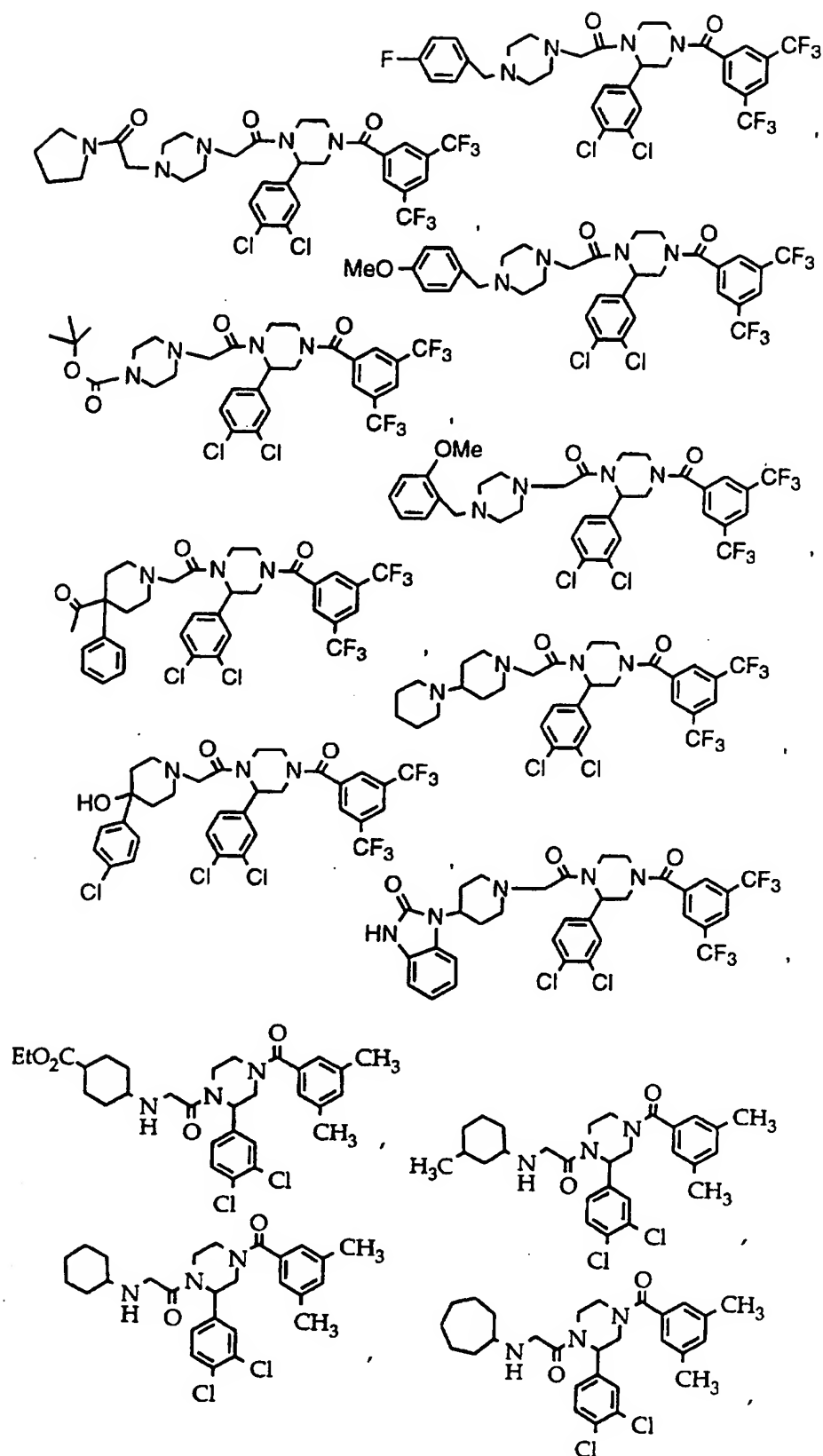


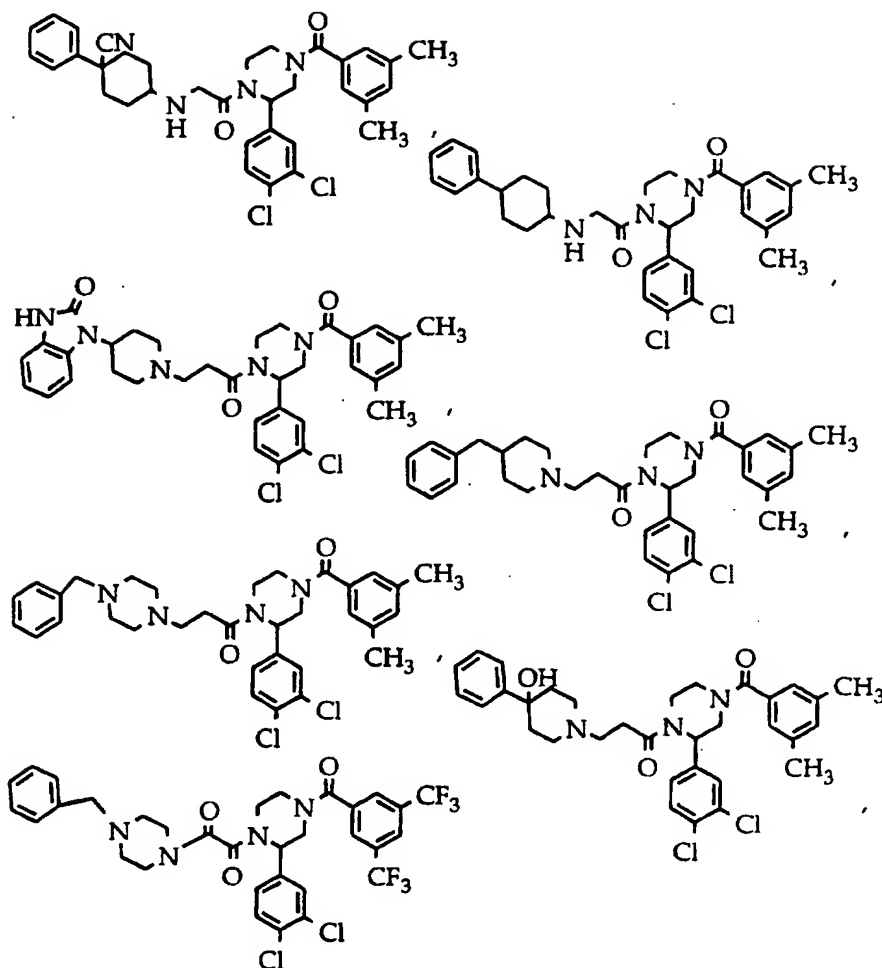
Where Z is



or a compound selected from the group consisting of







or a pharmaceutically acceptable salt thereof.

The invention also relates to a pharmaceutical composition comprising a therapeutically effective amount of a compound of formula I in combination with a pharmaceutically acceptable carrier.

The invention also relates to a method for inducing neurokinin antagonism which comprises administering a neurokinin antagonistic effective amount of a compound of formula I to a mammal in need thereof.

The invention also relates to a method for treating chronic airway diseases such as asthma and allergies; inflammatory diseases such as inflammatory bowel disease, psoriasis, fibrositis, osteoarthritis, and rheumatoid arthritis; migraine; central nervous system disorders such as depression, psychosis, dementia, and Alzheimer's disease; Down's syndrome; neuropathy; multiple sclerosis; ophthalmic disorders; conjunctivitis; auto immune disorders; graft rejection; systemic lupus erythematosus; GI disorders such as Crohn's disease and ulcerative

colitis; disorders of bladder function; circulatory disorders such as angina; Raynaud's disease; emesis, coughing and pain. In particular, the invention also relates to a method of treating asthma which comprises administering to a mammal in need of such treatment an anti-asthma
 5 effective amount of a compound of formula I for such purpose.

The invention also relates to the use of a compound of formula I for the preparation of a medicament useful for inducing neurokinin antagonism and in the treatment of the diseases cited above.

10 Detailed Description of the Invention

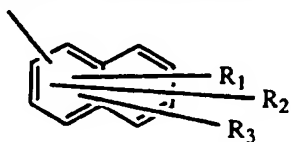
As used herein the term alkyl means a straight or branched, saturated hydrocarbon chain having from 1 to 6 carbon atoms. The number of carbon atoms may be designated. For example, "C₁-C₆ alkyl" represents a straight or branched, saturated hydrocarbon having from 1 to
 15 6 carbon atoms.

The term C₃-C₆ cycloalkyl means a cycloalkyl having from 3 to 6 carbon atoms, that is cyclopropyl, cyclobutyl, cyclopentyl and cyclohexyl.

The term alkenyl means means a straight or branched, saturated alkenyl having from 2 to 6 carbon atoms. The number of carbon atoms
 20 may be designated. For example, "C₂-C₆ alkenyl" represents a straight or branched alkenyl having from 1 to 6 carbon atoms.

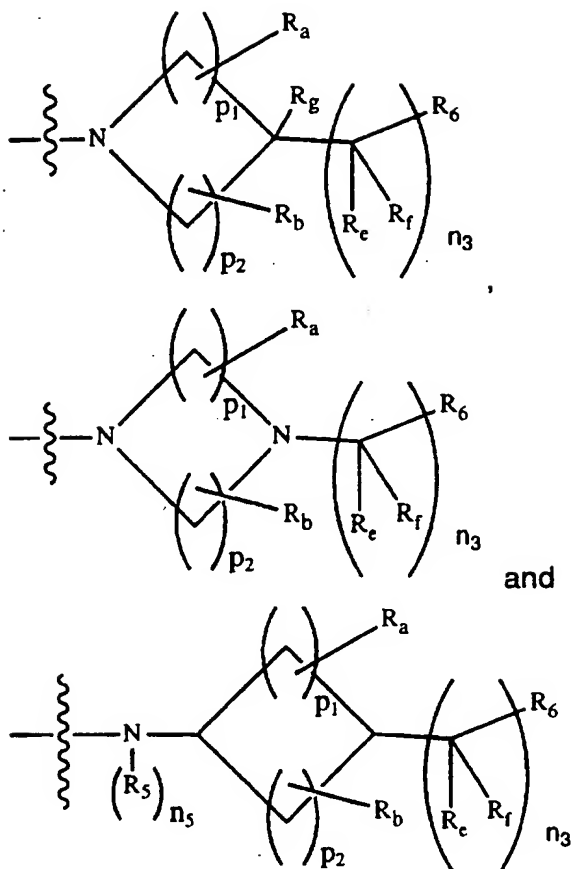
The term alkynyl means a straight or branched alkynyl having from 2 to 6 carbon atoms. The number of carbon atoms may be designated. For example, "C₂-C₆ alkynyl" represents a straight or branched chain alkynyl
 25 having from 2 to 6 carbon atoms.

As used herein, a heavy dark line (—) denotes a chemical bond coming above the plane of the page. A dashed line (·····) denotes a chemical bond coming below the plane of the page.

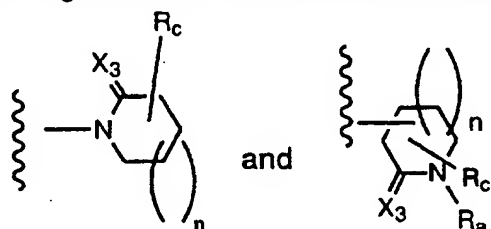


As used herein,
 30 R₁, R₂, and R₃ can be in either of the rings of the above naphthyl moiety.

In the rings shown below, one R_a and one R_b can be present at any position on the ring which will allow for a substitution:



Similarly, in the rings shown below, one R_C can be present at any
5 position on the ring which will allow for a substitution:



Asymmetric centers exist in compounds of formula I of the invention. Accordingly, compounds of formula I include stereoisomers.

All such isomeric forms and mixtures thereof are within the scope of the present invention. Unless otherwise indicated, the methods of preparation disclosed herein may result in product distributions which include all possible structural isomers, although it is understood that physiological response may vary according to stereochemical structure. The isomers may be separated by conventional means such as fractional crystallization, preparative plate or column chromatography on silica,

alumina, or reversed phase supports or HPLC (high performance liquid chromatography).

Enantiomers may be separated, where appropriate, by derivatization or salt formation with an optically pure reagent, followed by
 5 separation by one of the aforementioned methods. Alternatively, enantiomers may be separated by chromatography on a chiral support.

The compounds of formula I can exist in unsolvated as well as solvated forms, including hydrated forms, e.g. the hemihydrate. In general, the solvated forms, with pharmaceutically acceptable solvents such as
 10 water, ethanol, and the like are equivalent to the unsolvated forms for the purposes of the invention.

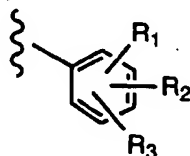
Those compounds of formula I which contain a basic group such as $-CH_2NH_2$, form pharmaceutically acceptable salts. The preferred pharmaceutically acceptable salts are nontoxic acid addition salts formed
 15 by adding to a suitable compound of the invention about a stoichiometric amount of a mineral acid, such as HCl, HBr, H_2SO_4 or H_3PO_4 or of an organic acid such as acetic, propionic, valeric, oleic, palmitic, stearic, lauric, benzoic, lactic, para-toluenesulfonic, methanesulfonic, citric, maleic, fumaric, succinic and the like, respectively.

20

General Methods of Preparation

The compounds of this invention may be prepared by one of the following general methods. As used herein RT means room temperature. Unless otherwise indicated, variables in the structural formulas below are
 25 as defined above. Starting materials and reagents used in the methods and examples below, are known or may be prepared according to known methods.

As used herein the term "substituted phenyl" means



30 wherein R_1 , R_2 , and R_3 are as described herein.

"Substituted" means substituted by R_1 , R_2 , and/or R_3 as described herein.

"Aryl" means phenyl, naphthyl, indenyl, tetrahydronaphthyl, indanyl, anthracenyl or fluorenyl.

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"Halogeno" refers to fluoro, chloro, bromo or iodo atoms.

"Heterocycloalkyl" refers to 4- to 6-membered rings comprising 1 to 3 heteroatoms independently selected from the group consisting of -O-, -S- and -N(R⁶)-, with the remaining ring members being carbon.

- 5 Examples of heterocycloalkyl rings are tetrahydrofuranyl, pyrrolidinyl, piperidinyl, morpholinyl, thiomorpholinyl and piperazinyl.

- "Heteroaryl" refers to 5- to 10-membered single or benzofused aromatic rings comprising 1 to 3 heteroatoms independently selected from the group consisting of -O-, -S- and -N=. Examples of single-ring
10 heteroaryl groups are pyridyl, isoxazolyl, oxadiazolyl, furanyl, pyrrolyl, thienyl, imidazolyl, pyrazolyl, tetrazolyl, thiazolyl, thiadiazolyl, pyrazinyl, pyrimidinyl, pyridazinyl and triazolyl. Examples of benzofused heteroaryl groups are quinolinyl, thianaphthenyl and benzofurazanyl. N-oxides of nitrogen-containing heteroaryl groups are also included. All positional
15 isomers are contemplated, e.g., 1-pyridyl, 2-pyridyl, 3-pyridyl and 4-pyridyl.

- Where R² and R³ substituents form a ring and additional heteroatoms are present, the rings do not include adjacent oxygen and/or sulfur atoms or three adjacent heteroatoms. Typical rings so formed are
20 morpholinyl, piperazinyl and piperidinyl.

As used herein, the term "BOC" means t-butoxycarbonyl.

As used herein, the term "Ph" means phenyl.

As used herein, the term "RT" means room temperature.

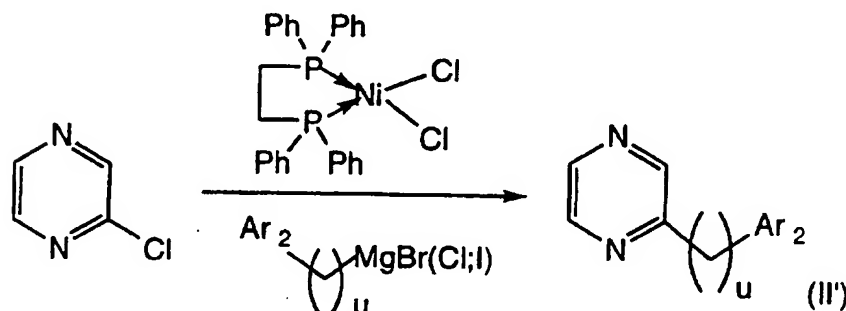
- As used herein, the term "parallel synthesis" means the preparation
25 of individual chemical compounds as one of a batch of, for instance, 20, 30, or even 100 identical reactions on usually a single substrate but using a different reagent in each vessel. Such reagents are always of the same general class- in this case, either carboxylic acids or organic amines in any set of parallel reactions. The conditions used for each reaction are
30 identical to those described in the examples, except that a simplified work-up is employed, generally a simple wash either with acid or base if appropriate, then water. The presence of the product is detected by thin layer chromatography (TLC) using known products as representative standards. Further characterization by combination HPLC/MS is generally
35 performed. No further purification is performed on these materials before they are submitted to biological assays.

As used herein, each R_C and R_C' is independently selected from the group consisting of H, C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, unsubstituted or substituted phenyl, and unsubstituted or substituted benzyl,

- 5 The starting materials in the methods below are either known or can be prepared in accordance with known methods. In particular, the following compounds are either known or can be prepared in accordance with known methods: the diamine A, the compounds of formulas A, VI, VIII, X, XI, XIV, XVIII, XIX, XXa, A', XXV, and Z-H, as well as esters of

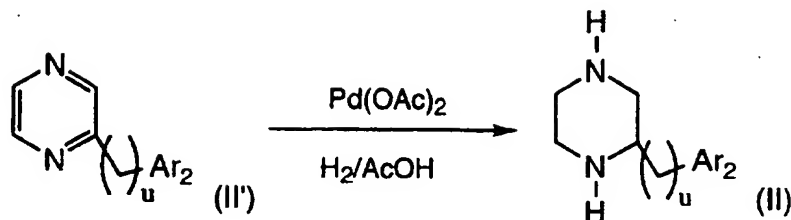
- 10 formula XI, and compounds of formula $Ar_2-(CH_2)_{u+1}-COO\text{-alkyl}$

Method 1. If the group Ar_2 is an aromatic group with no I or Br substituents, then the following method may be used to prepare the useful intermediates (IV):



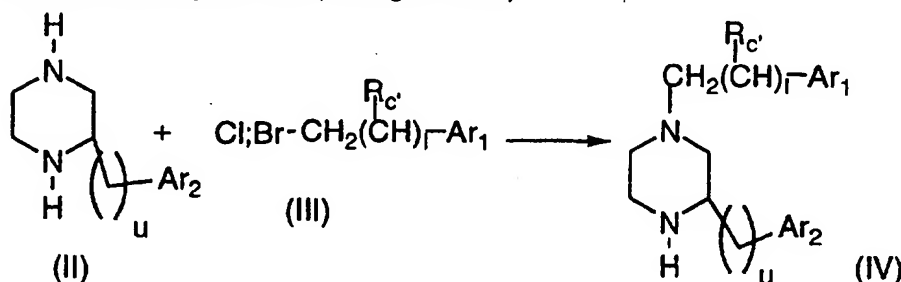
- 15 Transition metal catalyzed coupling of 2-chloropyrazine with an aromatic Grignard reagent in a dry, ether solvent, such as THF, yields the aryl-substituted pyrazine of formula II'. The catalyst shown, [1,2-bis-(diphenylphosphino)ethane]nickel(II) chloride, is a preferred reagent for this transformation. Where Ar_2 has no halo substituents, reduction of a
- 20 compound of formula II' by catalytic hydrogenation, using, for instance, palladium acetate, preferably in acetic acid solvent, results in preferential reduction of the pyrazine ring, leaving the aromatic ring unreduced, that is, it results in a compound of formula II. Similarly, 10% Pd on charcoal (Pd-C) can be used in an alcohol solvent, preferably methanol, with or without
- 25 the addition of a small quantity (1 to 5 equivalents) of acetic acid. Reaction times of about 1 to 24 hours generally suffice for this reaction, which is preferentially run at room temperature or slightly above (up to about 50°C) and using from 1 to about 6 atmospheres pressure of hydrogen.

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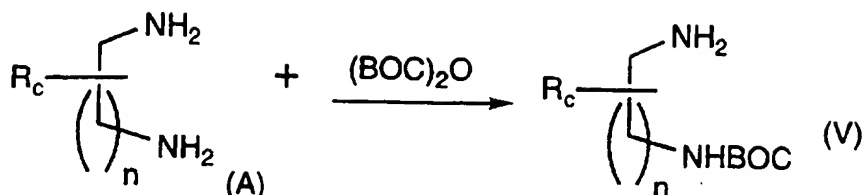


The intermediate of formula II may also be prepared from a compound of formula II', even if the group Ar₂ contains halogen atoms, by reduction using a strong hydride ion donor, preferably lithium aluminum hydride (LAH) or diisobutyl aluminum hydride (DIBAL-H) in an ether solvent, such as ether, THF or dimethoxyethane (DME).

Selective alkylation of a compound of formula II is possible using low temperature conditions. Thus, reacting a compound of formula II with a substituted aryl-alkyl halide of formula III where I is 0 to 2, results in the formation of the 4-substituted derivative of formula IV. Suitable conditions include use of a halogenated solvent, such as CH₂Cl₂, at low temperature. Suitable temperatures are from -78°C initially, allowing the reaction mixture to warm gradually to RT if the reaction is not completed after several hours. The reaction is catalyzed by the addition of an equivalent amount of an organic base, such as triethylamine and diisopropylethylamine (Hünig's base).

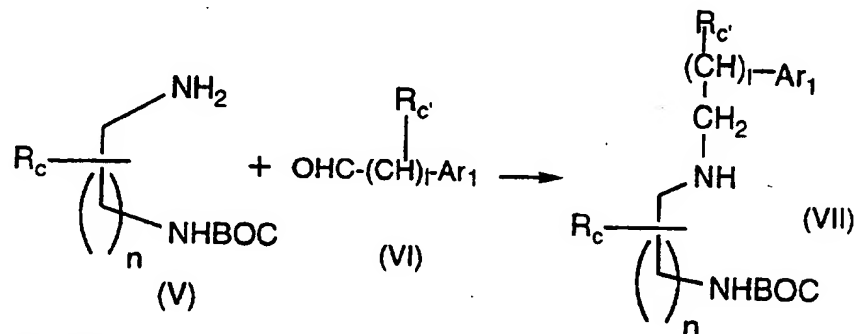


Method 2. If the group Ar₂ contains one or more halogen atoms on an aromatic ring and the other groups are as in Method 1, then an alternate route to a compound of formula IV is preferred. In addition, this method can be used to prepare compounds in which I is from 0 to 2. Mono-protection of the diamine of formula (A), preferably with BOC anhydride, or other agents known to introduce the t-butyloxycarbonyl protecting group, in an alcohol solvent, such as methanol, preferably at about -10°C, produces a compound of formula V.

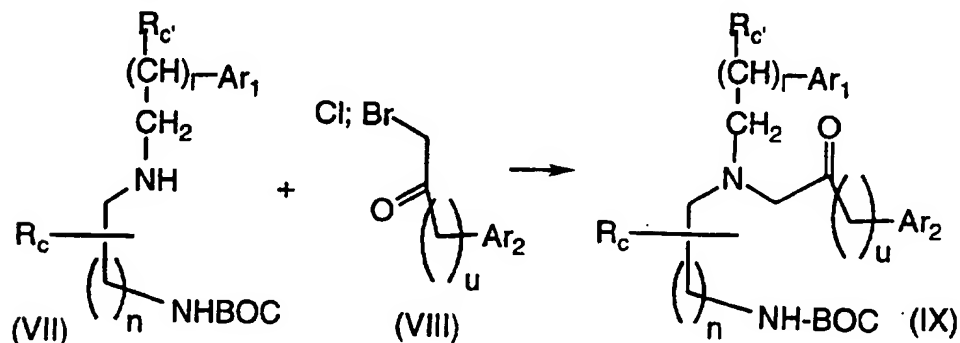


These compounds are used to perform a reductive amination reaction with the aldehyde of formula VI to produce an amine of formula VII. (In structures (A), (V), (VII), and (IX) herein, R_C can be bound to any position between the two nitrogens. In cyclic structures like (IVA) below, R_C can be bound to any available cyclic position that is occupied by carbon, and that is between the two nitrogens.)

Suitable conditions for this type of reaction include the use of an alcohol solvent, preferably methanol, or 2,2,2-trifluoroethanol, made slightly acidic with a weak organic acid, such as acetic acid, and a reducing agent known to favor reductive amination reactions, preferably sodium cyanoborohydride, NaBH_3CN .

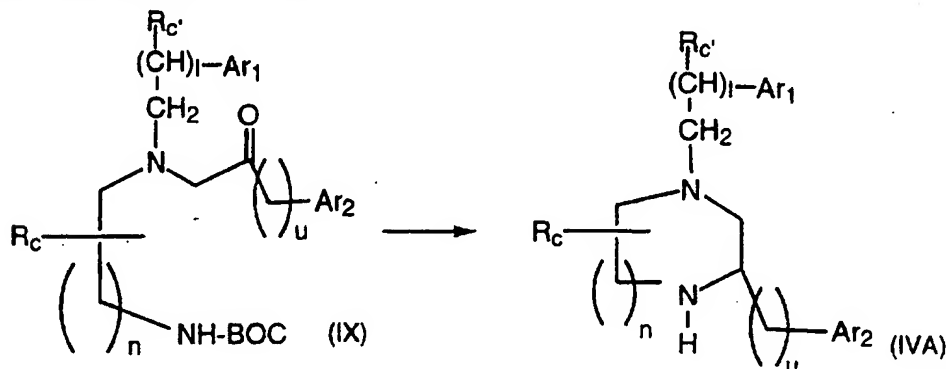


15 Reaction of a compound of formula VII with an α -haloketone of formula VIII, in which Ar₂ preferably represents a halogenated aromatic ring, but may be any of the claimed aromatic rings, in the presence of an organic base, such as di-isopropylethylamine, also known as Hünig's Base, in an ether solvent, such as THF, results in the formation of the intermediates of formula IX.



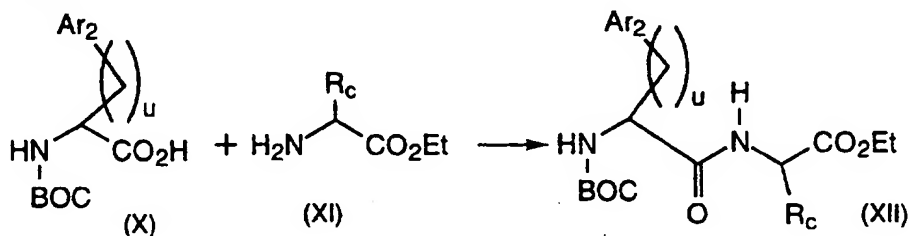
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Removal of the BOC protecting group using a suitable acidic catalyst, such as trifluoroacetic acid, followed by an intramolecular reductive amination, under conditions such as those described above for the preparation of a compound of formula VII, leads to the formation of compounds of formula IVA.



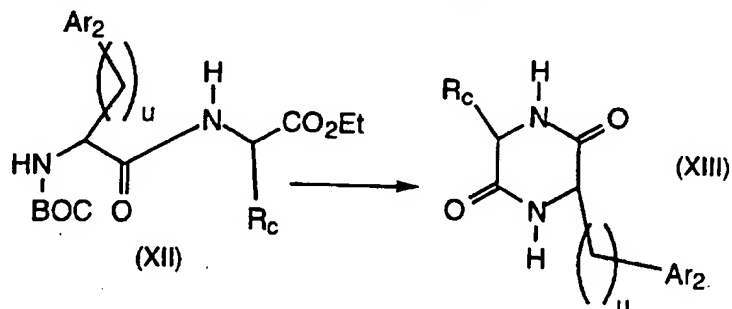
Method 3. An alternate route to compounds of the invention in which I is 0 to 2 is as follows. Standard coupling of an N-protected amino acid of formula X, wherein Ar₂ is as described above, with an amino acid ester

derivative $\text{H}_2\text{N}-\text{CH}(\text{R}_c)-\text{COOR}'$ (R' is C₂-C₄ alkyl, preferably, the ethyl ester of formula XI; Et in the formulas herein means ethyl), produces a dipeptide of formula XII. A suitable protecting group is BOC, although many others may also be used. Other esters of the amino acid may also be used. Standard coupling techniques may be applied, an example being the use of N-hydroxybenztriazole (HOBT) and a water-soluble carbodiimide, such as 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide (DEC), in a non-hydroxylic solvent such as CH₂Cl₂, DMF or a mixture of the two foregoing solvents. The reaction is run, preferably, at or below RT, and takes from 1 to 40 hours for completion, depending upon the substrates.

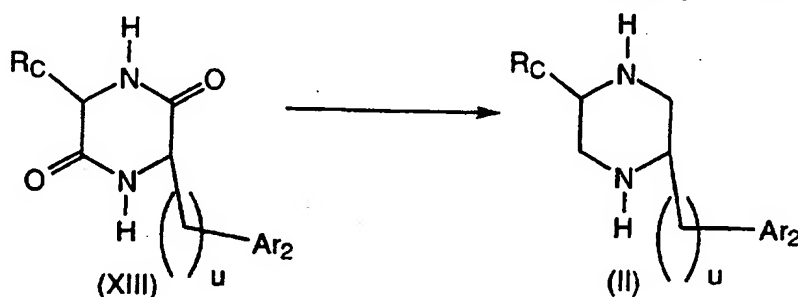


Removal of the protecting group under standard conditions, followed by treatment of the product with a base results in cyclization to

the diketopiperazine of formula XIII. Suitable conditions for removal of the exemplified BOC group are well known in the art and include catalysis by trifluoroacetic acid (TFA). A suitable base for cyclization is the alkali metal salt of an alcohol in the alcohol itself used as solvent. For example, a solution of sodium ethoxide in ethanol may be used. The temperature is preferably around RT but may be slightly above or below, in the range 0°C to about 40°C. The reaction is generally complete within a few hours. Suitable reaction times are from 1 to 24 hours.

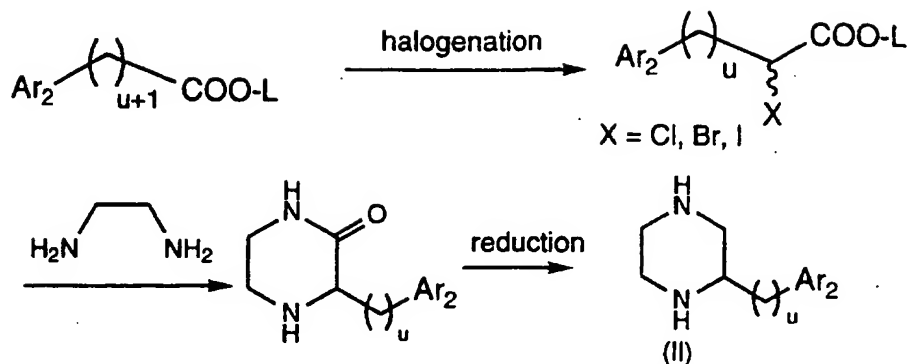


Reduction of the diketopiperazine of formula XIII to a compound of formula II may be accomplished preferentially with a strong hydride reducing agent, such as LAH or a solution of sodium bis(2-methoxyethoxy)aluminum hydride in toluene (also known as Red-Al®), or the $\text{BH}_3 \cdot \text{S}(\text{CH}_3)_2$ complex. Suitable solvents for this reaction are DME and other higher boiling ethers since the reaction is run at elevated temperatures, from about 50°C to about 110°C, preferably at about 90°C.



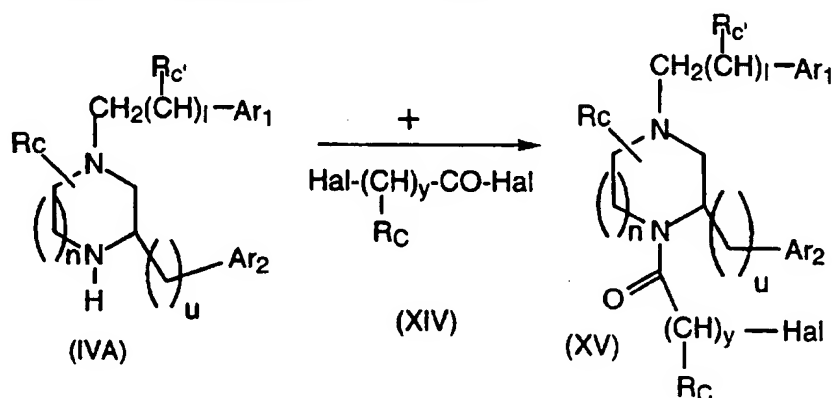
Alternatively, a compound of formula of II may be prepared by the scheme shown below (J. Med. Chem., 9, 181 (1966)). As used herein L is any readily available ester residue such as C_1 - C_7 alkyl, more preferably methyl or ethyl.

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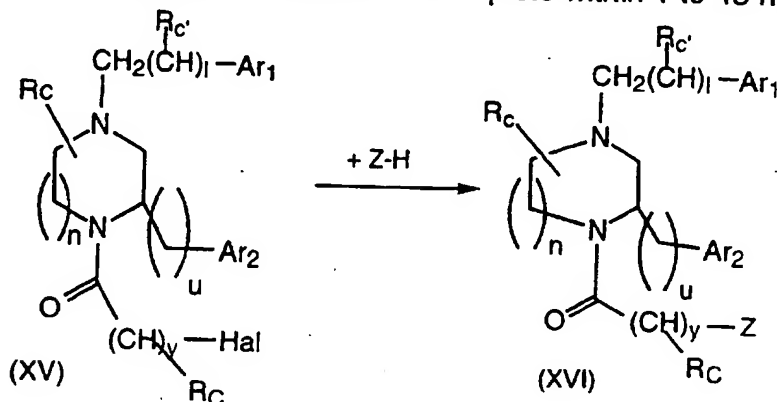
A compound of formula II may be converted to a compound of formula IV by the processes described in Method 1 above or Method 6 below.

- 5 **Method 4.** The intermediates of formula IV or IVA, formed via any of the previous methods, may be further processed as follows. A compound of formula IVA will be used in the Schemes. Reaction of a compound of formula IVA with an activated halo-acid, generally the acid halide of formula XIV, in which Hal represents Cl, Br, or I, yields the acylated derivative of formula XV that is, m is 1 for formula I. An organic base is used to take up the hydrogen halide formed in the reaction, suitable bases being triethylamine (TEA) and Hünig's Base. Suitable reaction media include halogenated solvents, such as methylene chloride and chloroform. The reaction is preferably run at low temperature, at least initially. Suitable temperatures are in the region of -50°C down to -80°C. Later in the reaction it may be desirable to allow the mixture to warm up to about RT to ensure completion of the reaction.



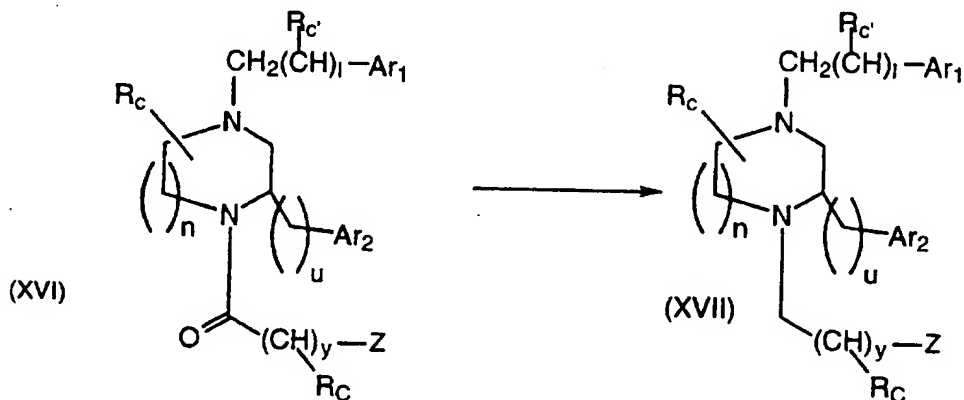
- 20 Reaction of the halogenated amides of formula XV with an amine of formula Z-H results in formation of the products of formula XVI, which are compounds of the invention in which X is O and m is 1. Compounds of formula XVI have been modified to show the fact that these products

could have been prepared from compounds of formula IVA as well as from IV. Suitable solvents for this reaction are halogenated hydrocarbons, such as methylene chloride, and an organic base is present to absorb the H-Hal formed. Appropriate bases include Hünig's Base. The reaction is performed at or around RT, a suitable temperature being generally in the range of from 0°C to 40°C. Reaction is complete within 1 to 48 hours.



Method 5.

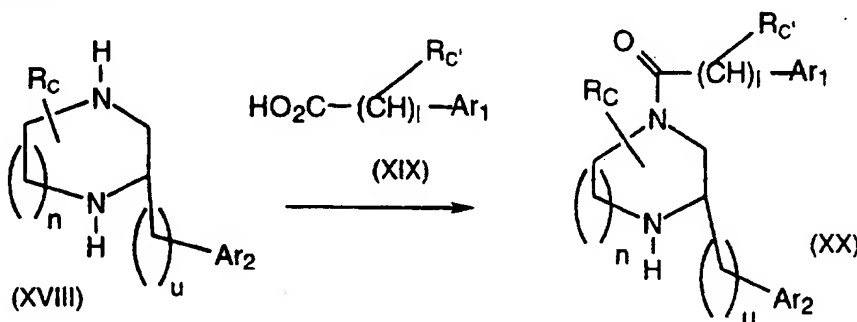
Compounds of formula XVI where $y \neq 0$ may be converted to other compounds of the invention of formula XVII by reduction under controlled conditions.



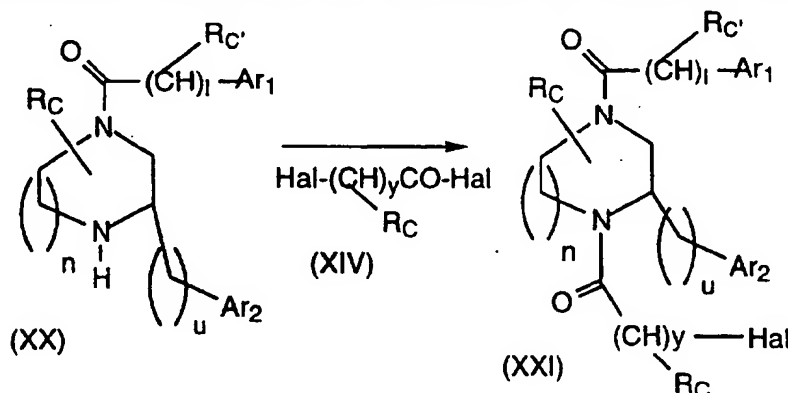
Suitable reducing agents to effect this transformation include the borane-dimethyl sulfide complex, as well as other less selective reagents, such as LAH, (assuming that no other group reactive to LAH is present), Red-Al®, and diborane in ether. Effective temperatures for the borane-dimethylsulfide complex to reduce compounds of formula XVI, range from RT to the reflux temperature of the solution of the reagent in THF (about 80°C).

Method 6. Intermediates of the formula XVIII may be selectively acylated by coupling with an acid of the formula XIX. Standard coupling

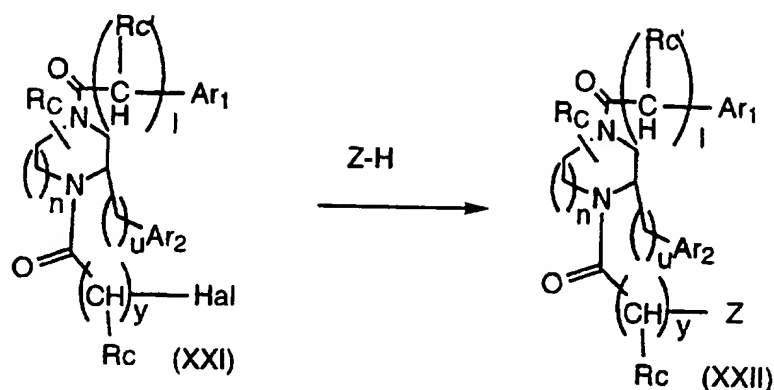
techniques may be applied, an example being the use of HOBT, a water-soluble carbodiimide, such as DEC, and an organic base, such as triethylamine, in a non-hydroxylic solvent, such as CH_2Cl_2 , at a temperature of about -20°C initially. The mixture may be allowed to warm to RT to complete the reaction. The product of reaction is the amide of formula XX.



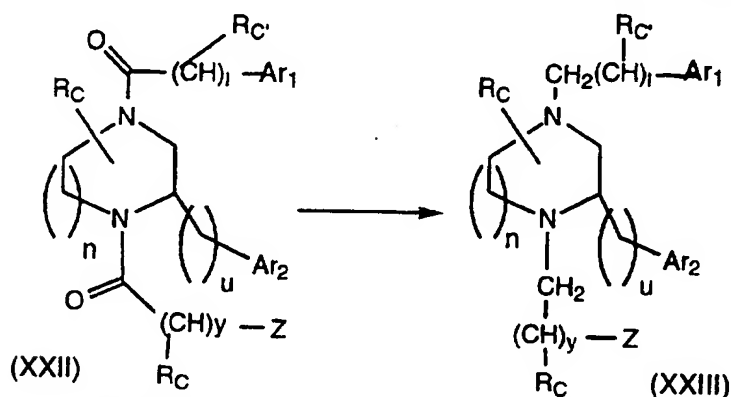
Compounds of the formula XX, may be further acylated using an acid halide of formula XIV. The reaction is run, preferably at about -78°C , over a period of 1 to 12 hours, in a halogenated solvent, such as methylene chloride or similar solvent. An organic tertiary amine is used to absorb the H-Hal produced in the reaction. Suitable amines include triethylamine and Hünig's Base. As used herein Hal means Cl, Br, or I.



The compounds of formula XXI, that is, m is 1 in formula I, $y = 1-3$, $l = 0-2$ may be used for further reaction without isolation. Additional organic base, for instance, Hünig's Base, is added to the mixture followed by Z-H, at or around -78°C . The reaction is completed by allowing the mixture to warm to RT overnight yielding the compounds of formula XXII after work-up and purification by standard methods.



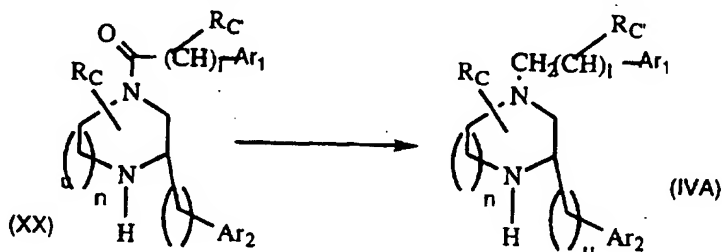
The compounds of formula XXII, in which $y = 1-3$, may be converted to other products of formula XXIII by reduction under controlled conditions.



- 5 Suitable reducing agents to effect this transformation include the borane-methyl sulfide complex, as well as other less selective reagents, such as LAH, Red-Al®, and diborane in ether or other unreactive solvents, such as THF. Using the borane-methyl sulfide complex in THF, at the reflux temperature of the solution, which is about 80°C, the reaction is
- 10 complete in about 2 hours to 48 hours depending on the precise substrate.

The Z-H substrates are commercially available or can be made by processes known to those skilled in the art. Specific examples are made according to the detailed procedures disclosed below.

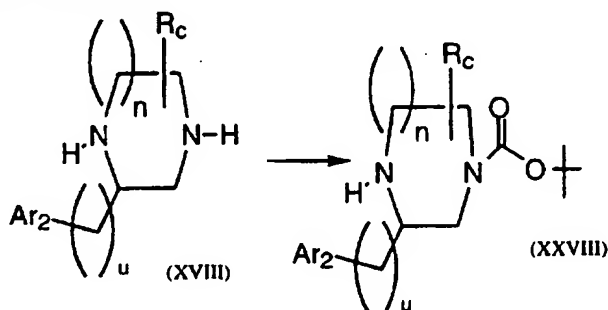
- 15 **Method 7.** The acylated derivatives of formula XX from Method 6 may be reduced to the saturated alkyl chain derivatives of formula IVA.



The process to conduct this conversion is the same as described in Method 6 for conversion of a compound of formula XXII to a compound of formula XXIII. The reagent of preference is the borane-methyl sulfide complex.

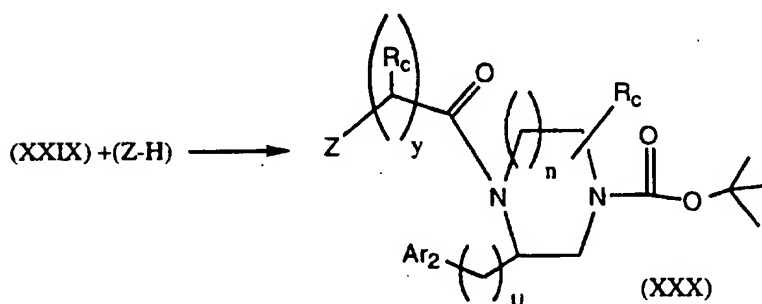
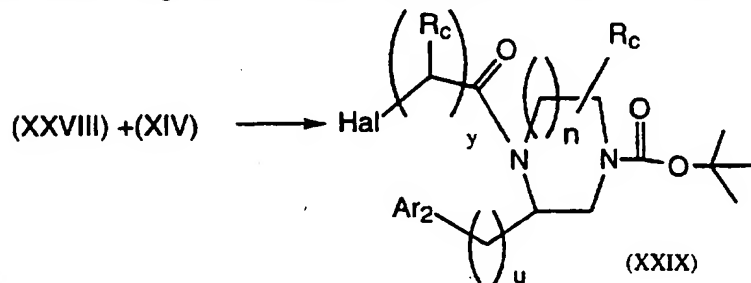
- 5 A compound of formula IVA can be converted to a target compound of formula XVI as described previously.

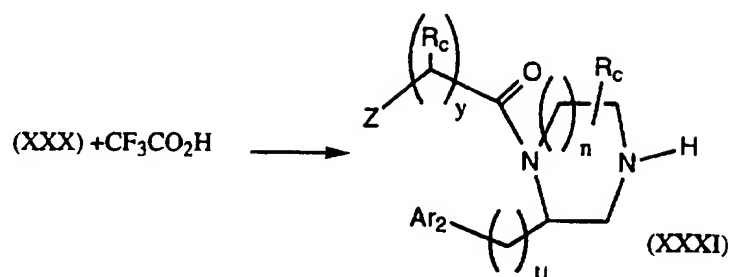
- 10 An alternate route to compounds of structure (XXII) also starts with compound (XVIII). Initial reaction with an amine protecting group reagent, preferably BOC anhydride, produces the N-t-butyloxycarbonyl derivative of the formula XXVIII.



As before, reaction occurs preferentially at the nitrogen atom further away from the Ar₂ group. Reaction of this intermediate with a reagent of structure (XIV) as described above, leads to the halo-derivative (XXIX).

- 15 Reaction of (XXIX) with Z-H, again as described above, produces the intermediate (XXX) which may be de-protected to produce (XXXI). Suitable reagents include trifluoroacetic acid and HCl.



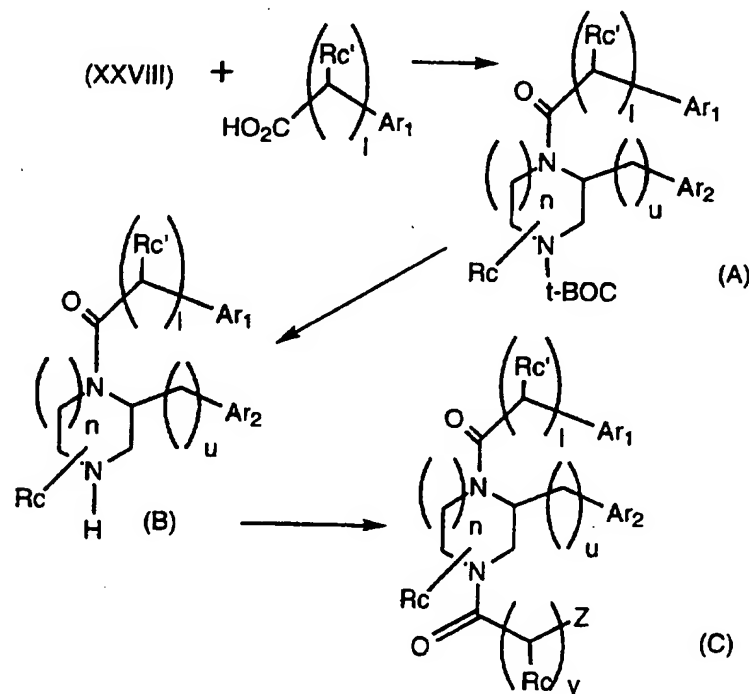


Reaction of (XXXI) with a carboxylic acid (XIX) under such coupling conditions as described above, leads to the products of formula (XXII).

- Method 7a.** Synthesis of the compounds of the invention wherein the pendant aromatic group Ar₂, or the pendant aromatic group Ar₂ and its sidechain, are located in the alternate ring position to the compounds of formula XXII (i.e. compounds of formula C below), may be prepared using compounds of formula XXVIII from method 7 as starting materials.

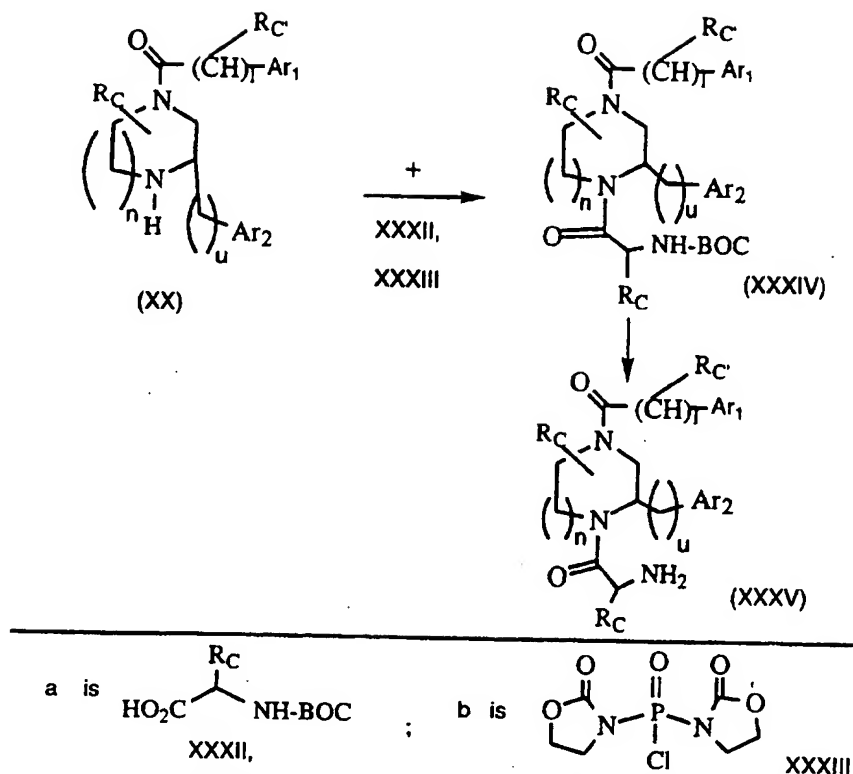
Coupling of compounds of formula XXVIII with any of the acids

- 10 $\text{Ar}_1-\overset{\text{Rc}'}{\underset{|}{\text{CH}}}_l\text{CO}_2\text{H}$ under standard coupling conditions, for instance using HOBT, Et₃N and DEC in CH₂Cl₂, produces the intermediate (A). Removal of the t-BOC or other protecting group under standard conditions releases the free amine (B). Acylation of (B) and further reaction with Z-H proceeds as described in Method 6 for the conversion of (XX) via (XXI) to (XXII) to
- 15 produce compound (C) of the invention.



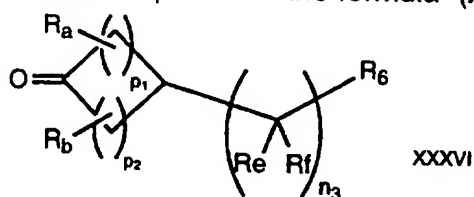
Method 8. A method for introducing a group, R_c , into the sidechain of a compound of the invention begins with a previously prepared compound of formula (XX). This may be coupled with a suitably protected amino-acid derivative of formula (XXXII) in which the t-BOC group is used as a representative protecting group. Use of a relatively reactive coupling agent, such as BOP-Cl of formula (XXXIII), is preferred and the reaction is run under standard coupling conditions well known to one skilled in the art. Suitable conditions include the use of CH_2Cl_2 and/or DMF as solvent, with triethylamine or Hünig's Base, and a temperature between 0°C initially and RT. Usual work-up conditions yield the protected intermediate of formula (XXXIV).

In the case of (XXXIV), in which the N-protecting group is t-BOC, the usual conditions for removal of such a group may be used to free the amine function. Various concentrations of CF_3CO_2H in CH_2Cl_2 will usually suffice. In some substrates a fairly dilute solution (e.g. 2N) will be sufficient whereas in other cases a more concentrated solution, up to neat TFA, may be necessary. In addition, other N-protecting groups may be employed and removed by methods well known in the art. An example is use of the N-Cbz which may be removed under either acidic or hydrogenolytic conditions. The result of deprotection is the amine intermediate of the formula (XXXV).

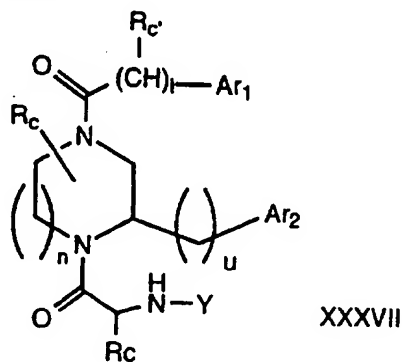


Conversion of intermediate of the formula (XXXV) to compounds of the invention is then carried out by a reductive alkylation process.

- 5 The group Z is introduced into the molecule using a ketone that is to be joined to the amino group of the formula (XXXV). An example of such an intermediate is a compound of the formula (XXXVI).



- 10 After the reaction this group becomes the Z group of the compounds of the invention, that is, the "Y-NH" group shown in compounds of the formula (XXXVII) just below



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is equivalent to the "Z" group shown in the Summary of the Invention. Conditions for this reductive amination procedure are known in the art and are exemplified by the use of NaBH_3CN in MeOH with the addition of several equivalents of acetic acid. Generally, the reaction is performed at RT and is left to react overnight.

Product is isolated by standard means, such as decomposition of excess reagent with H_2O and extraction of the product into an organic solvent such as CH_2Cl_2 or a mixture of Et_2O and CH_2Cl_2 .

Using procedures similar to those described in the above or using procedures known to those skilled in the art, one can produce all of the compounds of formula I of the invention. For example, one can obtain compounds of the invention of formula I wherein the R_c moiety is on various carbons of the piperazine ring.

The *in vitro* and *in vivo* activity of the compounds of formula I can be determined by the following procedures.

In vitro procedure to identify NK_1 activity

Test compounds are evaluated for their ability to inhibit the activity of the NK_1 agonist Substance P on the isolated guinea pig vas deferens. Freshly cut vas deferens are removed from male Hartley guinea pigs (230-350g) and suspended in 25 ml tissue baths containing Krebs' Henseleit solution warmed to 37°C and constantly aerated with 95% O_2 and 5% CO_2 . Tissues are adjusted to 0.5 g and allowed to equilibrate for a period of 30 minutes. The vas deferens are exposed to an electrical field stimulation (Grass S48 Stimulator) every 60 seconds at an intensity that will cause the tissue to contract 80% of its maximum capacity. All responses are recorded isometrically by means of a Grass force displacement transducer (FT03) and Harvard electronic recorder. Substance P inhibits the electrical field stimulated-induced contractions of the guinea pig vas deferens. In unpaired studies, all tissues (control or drug treated) are exposed to cumulative concentrations of Substance P ($1 \times 10^{-10} \text{ M}$ - $7 \times 10^{-7} \text{ M}$). Single log-concentrations of the test compounds are given to separate tissues and allowed to equilibrate for 30 minutes before a Substance P concentration-response curve is generated. At least 5 separate tissues are used for each control and individual drug-concentration for every drug assay.

Inhibition of the Substance P is demonstrated by a rightward shift of its concentration-response curve. These shifts are used to determine the

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pA₂ value, which is defined as the negative log of the molar concentration of the inhibitor which would require that twice as much agonist be used to elicit a chosen response. This value is used to determine relative antagonist potency.

5 Isolated Hamster Trachea NK₂ Assay

General methodology and characterization of hamster trachea responses to neurokinin agonists as providing an NK₂ monoreceptor assay is found in C.A. Maggi, et al., *Eur. J. Pharmacol.* 166 (1989) 435 and J.L. Ellis, et al., *J. Pharm. Exp. Ther.* 267 (1993) 95.

10 Continuous isometric tension monitoring is achieved with Grass FT-03 force displacement transducers connected to Buxco Electronics preamplifiers built into a Graphtec Linearcorder Model WR 3310. Male Charles River LAK:LVG (SYR) hamsters, 100-200 g fed weight, are stunned by a sharp blow to the head, loss of corneal reflex is assured, the
15 hamsters are sacrificed by thoractomy and cutting the heart. Cervical trachea segments are removed to room temperature Krebs buffer, pH 7.4, aerated with 95% O₂ - 5% CO₂ gas and cleaned of adhering tissue. The segments are cut into two 3-4 mm long ring segments. Tracheal rings are suspended from transducers and anchored in 15.0 ml water jacketed
20 organ baths by means of stainless steel hooks and 6-0 silk. Baths are filled with Krebs buffer, pH 7.4, maintained at 37°C and continuously aerated with 95% O₂ - 5% CO₂ gas. Tracheal rings are placed under 1.0 g initial tension and allowed a 90 min equilibration period with four 1 µM NKA challenge, wash and recovery cycles at 20 min intervals. 30 min
25 vehicle pretreatment is followed by cumulative additions of rising doses of NKA (3 nM - 1 µM final concentration, 5 min intervals between additions). The final NKA response is followed by a 15 min wash and recovery period. 30 min pretreatment with a test compound or its vehicle is followed by cumulative additions of rising doses of NKA (3 nM - 10 µM
30 final concentration if necessary, 5 minutes intervals between additions). The final NKA response is followed by a 1 mM carbachol challenge to obtain a maximal tension response in each tissue.

Tissue responses to NKA are recorded as positive pen
displacements over baseline and converted to grams tension by
35 comparison to standard weights. Responses are normalized as a % of the maximal tissue tension. ED₅₀'s are calculated for NKA from the control and treated NKA dose responses and compared. Test compounds

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resulting in an agonist dose ratio ≥ 2 at a screening concentration of 1 μM (i.e. $\text{pA}_2 \geq 6.0$) are considered actives. Further dose response data is obtained for actives so that an apparent pA_2 estimate can be calculated. pA_2 is calculated either by estimation of K_i as described by Furchgott (where $\text{pA}_2 = -\text{Log } K_i$, R.F. Furchgott, *Pharm. Rev.* 7 [1995] 183) or by Shild Plot Analysis (O. Arunlakshana & H.O. Shild, *Br. J. Pharmacol.* 14[1959] 48) if the data is sufficient.

Effect of NK₁ Antagonists on Substance P-Induced Airway Microvascular Leakage in Guinea Pigs

- 10 Studies are performed on male Hartley guinea pigs ranging in weight from 400-650 g. The animals are given food and water *ad libitum*. The animals are anesthetized by intraperitoneal injection of dialurethane (containing 0.1 g/ml diallylbarbituric acid, 0.4 g/ml ethylurea and 0.4 g/ml urethane). The trachea is cannulated just below the larynx and the
- 15 animals are ventilated ($V_T = 4$ ml, $f = 45$ breaths/min) with a Harvard rodent respirator. The jugular vein is cannulated for the injection of drugs.
- The Evans blue dye technique (Danko, G. et al., *Pharmacol. Commun.*, 1, 203-209, 1992) is used to measure airway microvascular leakage (AML). Evans blue (30 mg/kg) is injected intravenously, followed
- 20 1 min later by i.v. injection of substance P (10 $\mu\text{g/kg}$). Five min later, the thorax is opened and a blunt-ended 13-gauge needle passed into the aorta. An incision is made in the right atrium and blood is expelled by flushing 100 ml of saline through the aortic catheter. The lungs and trachea are removed en-bloc and the trachea and bronchi are then blotted dry with
- 25 filter paper and weighed. Evans blue is extracted by incubation of the tissue at 37°C for 18 hr in 2 ml of formamide in stoppered tubes. The absorbance of the formamide extracts of dye is measured at 620 nm. The amount of dye is calculated by interpolation from a standard curve of Evans blue in the range 0.5-10 $\mu\text{g/ml}$ in formamide. The dye concentration is
- 30 expressed as ng dye per mg tissue wet weight. Test compounds were suspended in cyclodextran vehicle and given i.v. 5 min before substance P.

Measurement of NK₂ Activity In Vivo

- 35 Male Hartley guinea pigs (400-500 gm) with ad lib. access to food and water are anesthetized with an intraperitoneal injection of 0.9 ml/kg dialurethane (containing 0.1 g/m diallylbarbituric acid, 0.4 g/ml ethylurea and 0.4 g/ml urethane). After induction of a surgical plane of anesthesia,

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tracheal, esophageal and jugular venous cannulae are implanted to facilitate mechanical respiration, measurement of esophageal pressure and administration of drugs, respectively.

5 The guinea pigs are placed inside a whole body plethysmograph and the catheters connected to outlet ports in the plethysmograph wall. Airflow is measured using a differential pressure transducer (Validyne, Northridge CA, model MP45-1, range ± 2 cmH₂O) which measures the pressure across a wire mesh screen that covers a 1 inch hole in the wall of the plethysmograph. The airflow signal is electrically integrated to a
10 signal proportional to volume. Transpulmonary pressure is measured as the pressure difference between the trachea and the esophagus using a differential pressure transducer (Validyne, Northridge, CA, model MP45-1, range ± 20 cm H₂O). The volume, airflow and transpulmonary pressure signals are monitored by means of a pulmonary analysis computer (Buxco
15 Electronics, Sharon, CT, model 6) and used for the derivation of pulmonary resistance (R_L) and dynamic lung compliance (C_{Dyn}).

Bronchoconstriction Due to NKA

Increasing iv doses of NKA are administered at half log (0.01-3
20 $\mu\text{g/kg}$) intervals allowing recovery to baseline pulmonary mechanics between each dose. Peak bronchoconstriction occurs within 30 seconds after each dose of agonist. The dose response is stopped when C_{Dyn} is reduced 80-90% from baseline. One dose-response to NKA is performed in each animal. Test compounds are suspended in cyclodextran vehicle and given i.v. 5 min before the initiation of the NKA dose response.

25 For each animal, dose response curves to NKA are constructed by plotting the percent increase in R_L or decrease in C_{Dyn} against log dose of agonist. The doses of NKA that increased R_L by 100% (R_L100) or decreased C_{Dyn} by 40% ($C_{Dyn}40$) from baseline values are obtained by log-linear interpolation of the dose response curves.

Neurokinin Receptor Binding Assay(s)

Chinese Hamster ovary (CHO) cells transfected with the coding regions for the human neurokinin 1 (NK₁) of the human neurokinin 2 (NK₂) receptors are grown in Dulbecco's minimal essential medium supplemented with 10% fetal calf serum, 0.1 mM non-essential amino
35 acids, 2 mM glutamine, 100units/ml of penicillin and streptomycin, and 0.8 mg of G418/ml at 37°C in a humidified atmosphere containing 5% CO₂.

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Cells are detached from T-175 flasks with a sterile solution containing 5mM EDTA in phosphate buffered saline. Cells are harvested by centrifugation and washed in RPMI media at 40°C for 5 minutes. The pellet is resuspended in Tris-HCl (pH7.4) containing 1 μ M phosphoramidon and 4 μ g/ml of chymostatin at a cell density of 30×10^6 cells/ml. The suspension is then homogenized in a Brinkman Polytron (setting 5) for 30-45 seconds. The homogenate is centrifuged at $800 \times g$ for 5 min at 4°C to collect unbroken cells and nuclei. The supernatant is centrifuged in a Sorvall RC5C at 19,000 rpm ($44,00 \times g$) for 30 min at 4°C. The pellet is resuspended, an aliquot is removed for a protein determination (BCA) and washed again. The resulting pellet is stored at -80°C.

To assay receptor binding, 50 μ l of [3 H]-Substance P (9-Sar, 11-Met [02]) (specific activity 41 Ci/mmol) (Dupont-NEN) (0.8 nM for the NK-1 assay) or [3 H]-Neurokinin A (specific activity 114 Ci/mmol) (Zenca) (1.0 nM for the NK-2 assay) is added to tubes containing buffer (50 mM Tris-HCl (pH 7.4) with 1 mM $MnCl_2$ and 0.2% Bovine Serum Albumin) and either DMSO or test compound. Binding is initiated by the addition of 100 μ l of membrane (10-20 μ g) containing the human NK-1 or NK-2 receptor in a final volume of 200 μ l. After 40 minutes at room temperature, the reaction is stopped by rapid filtration onto Whatman GF/C filters which have been presoaked in 0.3% polyethylenimine. Filters are washed 2 times with 3 ml of 50 mM Tris-HCl (pH7.4). Filters are added to 6 ml of Ready-Safe liquid scintillation cocktail and quantified by liquid scintillation spectrometry in a LKB 1219 RackBeta counter. Non-specific binding is determined by the addition of either 1 μ M of CP-99994 (NK₁) or 1 μ M SR-48968 (NK₂) (both synthesized by the chemistry department of Schering-Plough Research Institute). IC₅₀ values are determined from competition binding curves and Ki values are determined according to Cheng and Prusoff using the experimentally determined value of 0.8 nM for the NK₁ receptor and 2.4 nM for the NK₂ receptor.

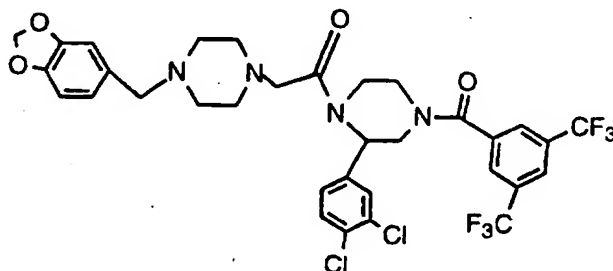
For all of the compounds of the invention, the NK₁ binding is in a range of about 0-100 % inhibition at 1 μ M concentration. For all of the compounds of the invention, the NK₂ binding is in a range of about 0-100 % inhibition at 1 μ M concentration. It should be understood that while the NK binding for certain compounds of the invention is as low as 0% at 1 μ M concentration, that at higher concentrations these compounds are expected to have NK binding inhibition activity.

The K_i of a compound is that concentration at which the compound caused 50% inhibition of either NK_1 or NK_2 . For those compounds of the invention having higher than 50% inhibition of NK_1 , K_i 's for NK_1 were determined. The K_i 's for NK_1 for such compounds fell within a range of about 0.1 nM to about 1 μ M.

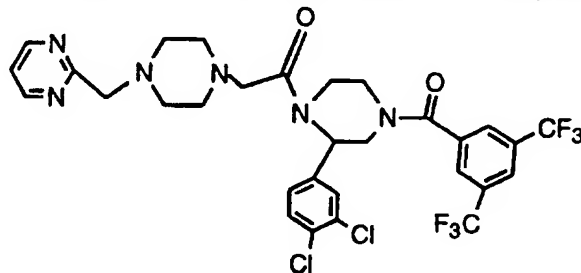
For those compounds of the invention having higher than 50% inhibition of NK_2 , K_i 's for NK_2 were determined. The K_i 's for NK_2 for such compounds fell within a range of about 0.1 nM to about 1 μ M.

Compounds of formula I exhibit NK_1 and NK_2 antagonist activity to varying degrees, i.e., certain compounds have strong NK_1 antagonist activity, but weaker NK_2 antagonist activity. Others are strong NK_2 antagonists, but weaker NK_1 antagonists. Certain compounds have both strong NK_1 and NK_2 antagonist activities. Some compounds can also be NK_3 antagonists.

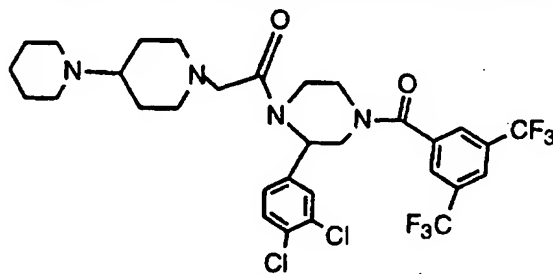
NK_1 binding and NK_2 binding values for certain compounds of the invention are as follows:



has a K_i for NK_1 binding of 5.3 nM; and has a K_i for NK_2 binding of 511 nM.

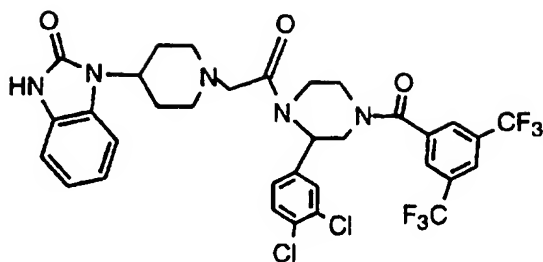


has a K_i for NK_1 binding of 23.3 nM; and a K_i for NK_2 binding of 29.1 nM.

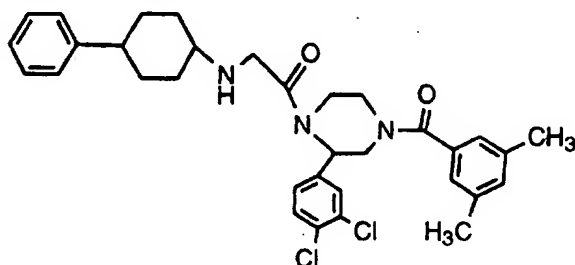


has a K_i for NK_1 binding of 3.3 nM; and a K_i for NK_2 binding of 93 nM.

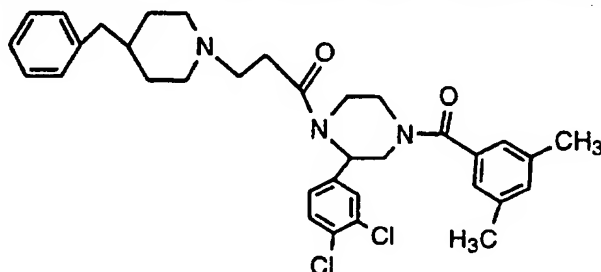
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has a K_i for NK_1 binding of 9.6 nM; and a K_i for NK_2 binding of 87 nM.

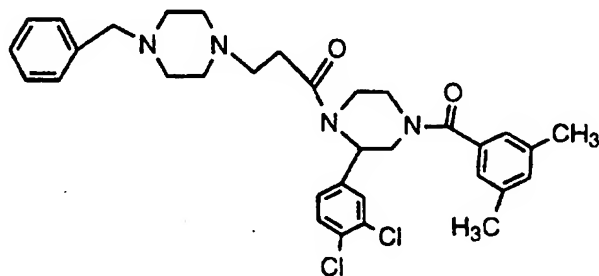


has a K_i for NK_1 binding of 7.0 nM; and a K_i for NK_2 binding of 40 nM.



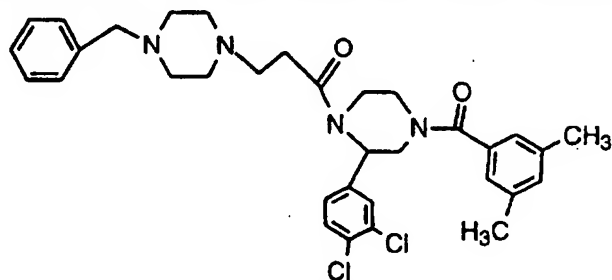
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has a K_i for NK_1 binding of 71 nM; and a K_i for NK_2 binding of 7.5 nM.



(Enantiomer B)

has a K_i for NK_1 binding of 67 nM; and a K_i for NK_2 binding of 18 nM.



10

(Racemic compound)

has a K_i for NK_1 binding of 3 nM; and a K_i for NK_2 binding of 25 nM.

Many compounds of formula I have an asymmetric center and therefore exist as a pair of enantiomers. In such cases, one enantiomer can have different biological activity than the other. For example, one enantiomer can have strong NK₁ activity and weak NK₂ activity while the other enantiomer has weak NK₁ activity and strong NK₂ activity.

Certain compounds of formula I have been found to be antagonists of both NK₁ and NK₂ receptors, and are therefore useful in treating conditions caused or aggravated by the activity of NK₁ and NK₂ receptors.

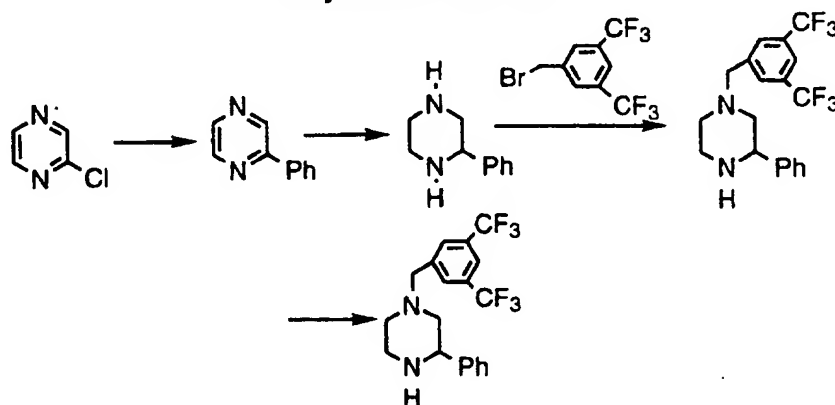
The present invention also relates to a pharmaceutical composition comprising a compound of formula I and a pharmaceutically acceptable carrier. Compounds of this invention can be administered in conventional oral dosage forms such as capsules, tablets, powders, cachets, suspensions or solutions, or in injectable dosage forms such as solutions, suspensions, or powders for reconstitution. The pharmaceutical compositions can be prepared with conventional excipients and additives, using well known formulation techniques. Pharmaceutically acceptable excipients and additives include nontoxic and chemically compatible fillers, binders, disintegrants, buffers, preservatives, anti-oxidants, lubricants, flavorings, thickeners, coloring agents, emulsifiers and the like.

The daily dose of a compound of formula I for treating asthma, cough, bronchospasm, inflammatory disease, migraine, nociception and gastrointestinal disorders is about 0.1 mg to about 20 mg/kg of body weight per day, preferably about 0.5 to about 15 mg/kg, more preferably 0.5 to about 5 mg/kg. For an average body weight of 70 kg, the dosage range is therefore from about 1 to about 1500 mg of drug per day, preferably about 50 to about 100 mg, given in a single dose or 2-4 divided doses. The exact dose, however is determined by the attending clinician, and is dependent on the potency of the compound administered, the age, weight, condition and response of the patient.

The invention disclosed herein is exemplified by the following examples, which should not be construed to limit the scope of the disclosure. Alternative mechanistic pathways and analogous structures within the scope of the invention will be apparent to those skilled in the art.

EXAMPLE 1

(+/-)-1-[[3,5-bis(trifluoromethyl)phenyl]methyl]-3-phenyl-piperazine,
dihydrochloride salt.



- 5 Chloropyrazine (20.68 g, 177 mmol) and [1,2-bis(diphenylphosphino)ethane]nickel(II) chloride (41.08 g, 77.8 mmol) in dry THF (1.5 L) were mixed and stirred for 80 minutes in a flask (cooled with a water bath) under nitrogen. A solution of phenylmagnesium bromide (3M in Et₂O) (103 mL, 309 mmol) was added slowly through a dropping funnel
- 10 into the cooled brick-red slurry at room temperature under nitrogen over 3.5 hours. After stirring at room temperature overnight, TLC showed that the reaction was complete. 3 N HCl (100 mL) was added slowly through a dropping funnel under nitrogen and the mixture was stirred for one hour. The THF layer was separated from the aqueous layer. The aqueous layer
- 15 was adjusted to pH 12 with 6 N NaOH and extracted with EtOAc (100 mL, 3x). The organic fractions (THF and EtOAc) were combined and dried over MgSO₄, filtered and concentrated to give a solid. The product was purified by flash chromatography on 300 g of flash grade silica gel in 2.5 % EtOAc/CH₂Cl₂ to give 10.79 gram (69 mmol, 39%) of 2-phenyl-
- 20 pyrazine, m.p. 69-70 °C; FAB MS [M+1]⁺ 157; Found, C, 76.55; H, 5.22; N, 17.71. Calcd. for C₁₀H₈N₂, C, 76.90; H, 5.16; N, 19.93.

To a solution of 2-phenylpyrazine (11.64 g, 74.53 mmol) in acetic acid (58.2 mL) was added palladium acetate Pd(OAc)₂ (2.33 g, 9.94 mmol). The mixture was hydrogenated at 50 psi for 4 h. After the reaction

25 was complete, the catalyst was filtered off and rinsed with a small portion of acetic acid. The filtrate was concentrated under house vacuum to give a brown-black solid which was suspended in deionized water (300 mL) and adjusted to pH 13 with 20 % NaOH solution. The product was extracted from aqueous solution with EtOAc (200 mL, 3x), dried over

MgSO₄, filtered and evaporated to dryness to give 2-phenylpiperazine (7.2 g). An additional 1.6 g of 2-phenylpiperazine was obtained by evaporating the aqueous fraction to a solid and triturating the solid with CH₂Cl₂. Total yield of 2-phenyl-piperazine was 73 %. The crude material

5 was crystallized from EtOAc and hexane for characterization, m.p. 86-88°C; FAB MS [M+1]⁺ 163; Found, C, 74.04; H, 8.66; N, 17.15. Calcd. for C₁₀H₁₄N₂, C, 74.04; H, 8.69; N, 17.26.

To a solution of 2-phenylpiperazine (4.0 g, 24.65 mmol) in dry CH₂Cl₂ (200 mL) at -78 °C under nitrogen was added Et₃N (5.15 mL, 36.97 mmol) followed by the dropwise addition of a CH₂Cl₂ solution

10 (46.60 mL) of bis(trifluoromethyl)benzyl bromide (4.66 mL, 24.65 mmol). The flask was kept at -78 °C then it was gradually warmed to room temperature overnight. After TLC showed that the reaction was complete, the material was washed with brine (150 mL, 2x), dried over MgSO₄,

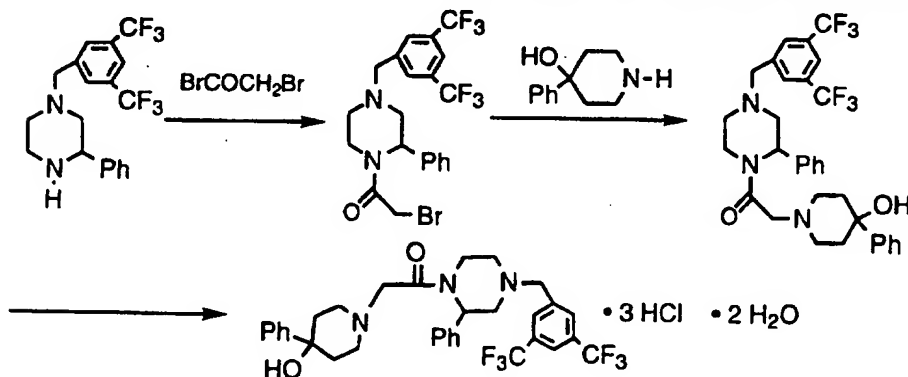
15 filtered, and evaporated under vacuum to yield a tan solid. The crude product was purified by flash silica gel chromatography (150 g), eluting with 2.5% MeOH/CH₂Cl₂ to give (+,-)-1-[[3,5-bis(trifluoromethyl)phenyl]-methyl]-3-phenylpiperazine (6.96 g, 17.92 mmol, 72.7%) as an oil. A

20 portion of this oil (0.5 g, 1.287 mmol) was converted to its hydrochloride salt by dissolving the oil in CH₂Cl₂ (20 mL) and treating with 2.3 M HCl-EtOH (1.3 mL 2.99 mmol). After stirring at room temperature for 10 minutes, all solvents were removed under high vacuum and the residue was dried overnight, m.p. 229-233 °C; FAB MS [M+1]⁺ 389;

25 Found, C, 48.83; H, 4.28; N, 5.87; Cl, 14.77; F, 24.03. Calcd. for C₁₉H₁₈N₂F₆•2 HCl•0.25 H₂O, C, 48.99; H, 4.43; N, 6.01; Cl, 15.22; F, 24.47.

EXAMPLE 2

(+,-)-4-[[3,5-bis(trifluoromethyl)phenyl]methyl]-1-[2-(4-hydroxy-4-phenyl-1-piperidiny)acetyl]-2-phenylpiperazine, dihydrochloride salt.



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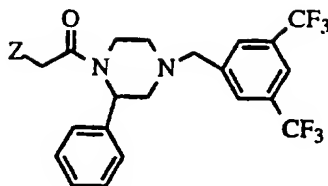
To a solution of (+,-)-1-[[3,5-bis(trifluoromethyl)phenyl]methyl]-3-phenylpiperazine (0.76 g, 1.975 mmol) in dry CH₂Cl₂ (15.2 mL) at -78°C was added Et₃N (0.286 mL, 2.055 mmol) followed by the dropwise addition of bromoacetyl bromide (0.179 mL, 2.055 mmol). After stirring at
5 -78 °C for 4 hours, the reaction was diluted with CH₂Cl₂ (200 mL), washed with brine (100 mL, 2x) and dried over MgSO₄. After filtration, the solvent was removed to give a light yellow solid which was used without further purification. FAB MS[M+1]⁺ 509.2 (⁷⁹Br).

The product from the previous reaction (2.0 g, 3.928 mmol) was
10 dissolved in dry CH₂Cl₂ (20 mL) and cooled to -78 °C under nitrogen. To this cooled solution were added 4-hydroxy-4-phenyl-piperidine (0.8 g, 4.49 mmol) and diisopropylethylamine (0.787 mL, 4.492 mmol). The reaction was gradually warmed to room temperature overnight under nitrogen. After completion, CH₂Cl₂ (300 mL) was added and the organic
15 layer was washed with brine (100 mL, 2x), dried over MgSO₄ and filtered. The filtrate was evaporated under vacuum to give a crude oil which was purified by flash chromatography on flash grade silica gel (100 g), eluting with 4.0% NH₃-MeOH-2.5% EtOH/CH₂Cl₂ to give a light yellow oil (2.24 g, 2.38 mmol, 94%). A portion of the oil (0.40 g, 0.661 mmol) was
20 converted to its hydrochloride salt by dissolving in CH₂Cl₂ (8 mL) and treating with 2.3 M HCl-EtOH (0.632 mL, 1.454 mmol). After stirring at room temperature for 30 minutes, solvent was evaporated and the residue was vacuum dried overnight, m.p. 185-187°C; FAB MS [M+1]⁺ 606.6 ; Found, C, 54.58; H, 5.44; N, 5.75; Cl, 9.71; F, 16.11. Calcd. for
25 C₃₂H₃₃O₂N₃F₆ • 2 HCl • 1.5 H₂O, C, 54.47; H, 5.43; N, 5.96; Cl, 10.05; F, 16.16.

EXAMPLE 3

By process analogous to that described in Example 2, employing appropriate heterocyclic derivatives (Z group), as listed below, in place of
30 4-hydroxy-4-phenylpiperidine, the following compounds were prepared.

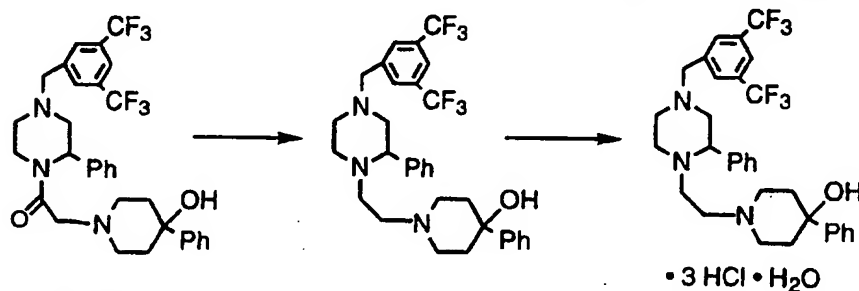
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Z	salt	m.p. °C	[M+1] FAB MS	High Res. MS
	free form	198-201	647.4	
	3 HCl • 2 H ₂ O	192-195	605.5	
	2 HCl • 1.5 H ₂ O	162-165	604.5	
	free form	66-68	619.3	Cal'd 619.2872 Found 619.2856
	free form	68-71	682.3	

EXAMPLE 4

(+,-)-1-[2-[4-[[3,5-bis(trifluoromethyl)phenyl]methyl]-2-phenyl-1-piperazinyl]ethyl]-4-phenyl-4-piperidinol, trihydrochloride salt.



5

To a solution of (+,-)-4-[[3,5-bis(trifluoromethyl)phenyl]methyl]-1-[2-(4-hydroxy-4-phenyl-1-piperidinyl)acetyl]-2-phenylpiperazine (0.74 g, 1.222 mmol) in THF (18.5 mL) was added 10 M BH₃ • S(CH₃)₂ (0.85 mL, 8.5 mmol). The mixture was heated in an oil bath at 80 °C under nitrogen overnight. After completion, excess BH₃ was decomposed by dropwise addition of MeOH to the cooled solution under nitrogen. MeOH was evaporated and the residue was redissolved in EtOH (22.2 mL). K₂CO₃ (0.31 g, 2.69 mmol) was added and the mixture was refluxed at 80 °C for five hours. After TLC showed that the reaction was complete, the solid

10

- 46 -

- was filtered off and the filtrate was evaporated under vacuum. The residue was redissolved in EtOAc (200 mL), washed with brine (100 mL) and dried over MgSO₄. It was filtered and evaporated under vacuum to give an oil which was purified by flash chromatography on flash grade silica gel (100 g), eluting with 10% NH₃-MeOH/CH₂Cl₂ to give the desired material as an oil (0.504 g, 0.852 mmol, 69.8 %). A portion of the oil (0.35 g) was converted to its hydrochloride salt by dissolving in dry CH₂Cl₂ (17.5 mL), followed by the addition of 2.3 M HCl-EtOH (0.84 mL). Solvents were removed after stirring at room temperature for 0.25 h and the residue was vacuum dried, m.p. 215-220 °C; FAB MS [M+1]⁺ 592.1; Found, C, 53.17; H, 5.51; N, 5.77; Cl, 14.37; F, 15.62. Calcd. for C₃₃H₃₈N₄F₆•3 HCl•H₂O, C, 53.45; H, 5.61; N, 5.84; Cl, 14.79; F, 15.85.

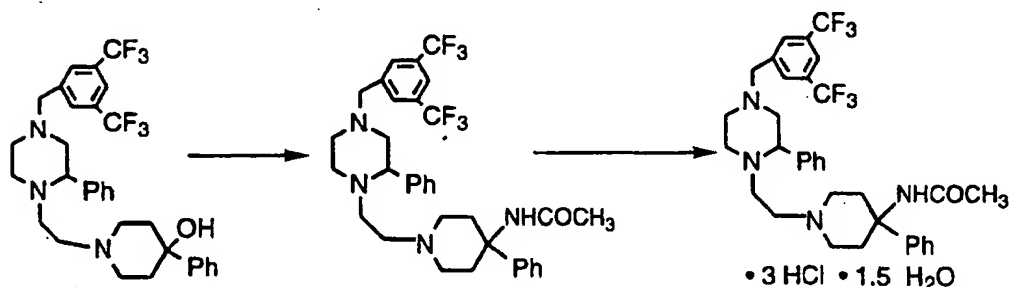
EXAMPLE 5

- By a process analogous to the process described in Examples 2 and 4, employing appropriate heterocyclic derivatives (Z group), listed below, in place of 4-hydroxy-4-phenylpiperidine, the following compounds were prepared.

Z	salt	m.p. °C	[M+1] ⁺ FAB MS
	4 HCl • 1.0 H ₂ O	> 230	591.1
	3 HCl • 1.0 H ₂ O	>260	590.9

EXAMPLE 6

- (+,-)-N-[1-[2-[4-[[3,4-bis(trifluoromethyl)phenyl]methyl]-2-phenyl-1-piperazinyl]ethyl]-4-phenyl-4-piperidiny]acetamide, trihydrochloride salt, 1.5 hydrate.



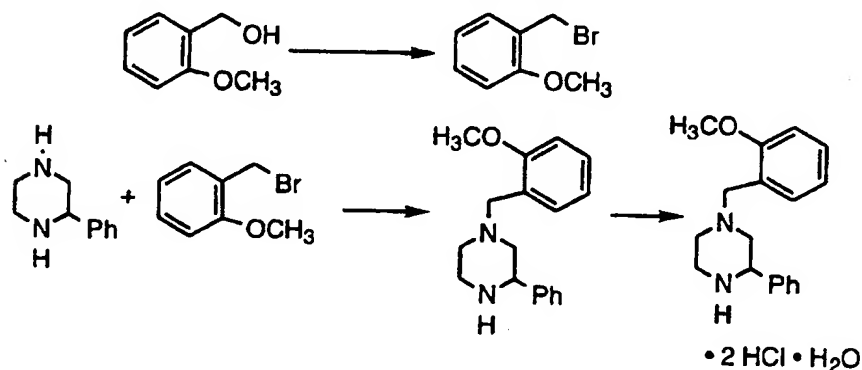
- 47 -

To a solution of the product from Example 4 (free form) (0.66 g, 1.116 mmol) in 5.0 ml of acetonitrile was added dropwise concentrated sulfuric acid (2.44 mL) at room temperature under nitrogen. After 4 hours, water (100 mL) was added to the reaction and the solution was adjusted to pH 9 with 10% NaOH solution. The product was extracted from the aqueous solution with EtOAc (100 mL, 3x). The organic fractions were combined and washed with brine (100 mL), dried over MgSO₄, filtered, and concentrated under vacuum to give an oil. The product was purified by flash chromatography on flash silica gel (80 g) and was eluted with 10% NH₃-MeOH/CH₂Cl₂ to give an oil (0.40 g, 0.774 mmol, 69%). A portion of this oil (0.4 g, 0.632 mmol) was dissolved in CH₂Cl₂ (20 ml) and treated with 2.3 M HCl-EtOH (2.528 mmol). After stirring at RT for 0.5 h, solvents were removed under vacuum to give a white solid, m.p. 245-247 °C; FAB MS [M+1]⁺ 633.4;

Found, C, 53.27; H, 5.80; N, 7.23; Cl, 13.91; F, 14.55. Calcd. for C₃₄H₃₈ON₄F₆ • 3 HCl • 1.5 H₂O, C, 53.09; H, 5.76; N, 7.28; Cl, 13.83; F, 14.82.

EXAMPLE 7

(+,-)-1-[(2-methoxyphenyl)methyl]-3-phenyl-piperazine, dihydrochloride salt.



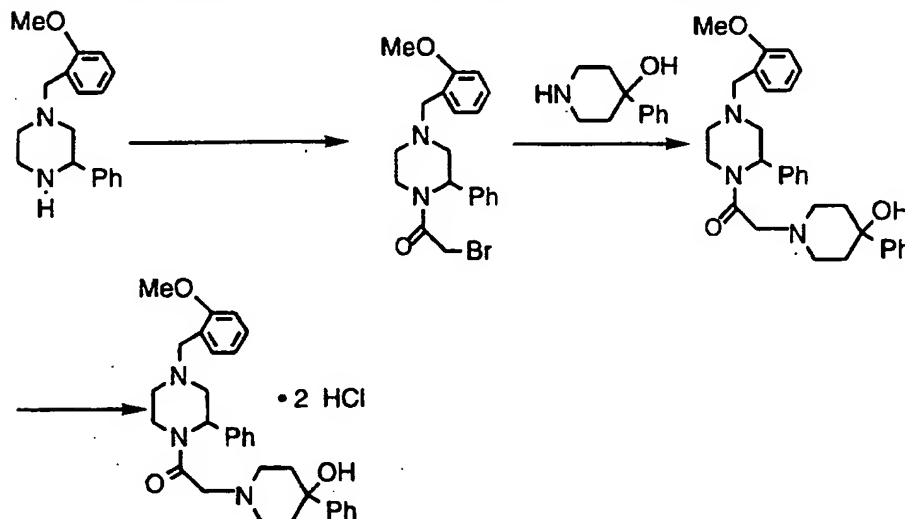
To a solution of 1-hydroxymethyl-2-methoxy-benzene (28.7g, 0.207 mol) in CH₂Cl₂ (574 mL) at 0 °C under nitrogen was added slowly PBr₃ (13.66 mL, 0.145 mol). After stirring for an additional 1.5 hours, MeOH (13.66 mL) was added and stirred for 5 minutes. To this mixture was added dropwise 10% Na₂CO₃ (2 mL) solution and stirred for 5 minutes. The mixture was then washed with 10% Na₂CO₃ (50 mL, 2x) and brine (100mL). It was dried over MgSO₄, and filtered. The filtrate was removed under vacuum to give an oil (40 g) of 1-bromomethyl-2-methoxy-benzene. This material was used without purification.

- 48 -

- To a solution of (+,-)-2-phenyl-piperazine (2.83 g, 17.44 mmol) (described in Example 1) in dry CH_2Cl_2 (141.5 mL) at -78°C was added slowly a solution of 1-bromomethyl-2-methoxy-benzene (3.507 g, 17.44 mmol) in dry CH_2Cl_2 (35 mL) under nitrogen. The reaction was stirred at
- 5 -78°C and gradually warmed to RT overnight. After completion, the product was diluted with CH_2Cl_2 (200 mL), washed with brine (100 mL), dried over MgSO_4 and filtered. The filtrate was removed under vacuum to give an oil. The product was purified by flash chromatography on flash grade silica gel (150 g), eluting with 4% $\text{MeOH}/\text{CH}_2\text{Cl}_2$ to give the title
- 10 compound as an oil (2.68 gram, 54%). A portion of this oil (0.33 g, 1.168 mmol) was dissolved in CH_2Cl_2 (10.0 mL) and treated with 2.3 M HCl (1.1 mL, 2.53 mol). After stirring at room temperature for 10 minutes, solvents were removed under vacuum to give a solid, m.p. $152-156^\circ\text{C}$; FAB MS $[\text{M}+1]^+$ 283.2.
- 15 Found, C, 58.18; H, 7.23; N, 7.33; Cl, 18.89. Calcd. for $\text{C}_{18}\text{H}_{22}\text{ON}_2 \cdot 2\text{HCl} \cdot \text{H}_2\text{O}$, C, 57.91; H, 7.02; N, 7.50; Cl, 18.99.

EXAMPLE 8

(+,-)-1-[(4-hydroxy-4-phenyl-1-piperidiny)acetyl]-4-[(2-methoxy-phenyl)methyl]-2-phenyl piperazine, dihydrochloride salt.



20

- The bromoacetyl derivative of the product from Example 7 was prepared according to the procedure described in Example 2. This intermediate was used for the next reaction without further purification. The title compound was prepared by employing a process analogous to
- 25 that described in Example 3, via the bromoacetyl derivative of (+,-)-1-[(2-

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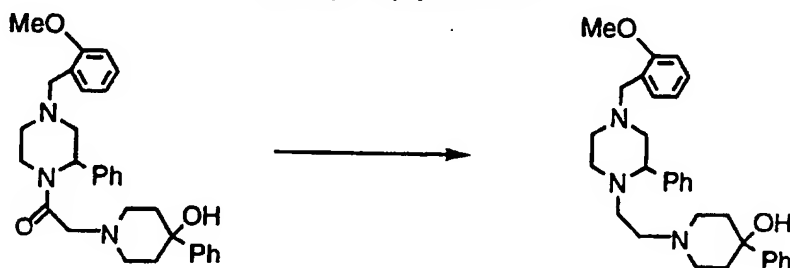
methoxyphenyl)methyl]-3-phenyl-piperazine to give a solid, m.p. 183-186 °C; FAB MS $[M+1]^+$ 500;

Found C, 61.30; H, 7.54; N, 6.98; Cl, 11.65. Calcd. for $C_{31}H_{37}N_3O_3 \cdot 2$ HCl, C, 61.18; H, 7.12; N, 6.90; Cl, 11.65.

5

EXAMPLE 9

(+,-)-1-[2-[4-[(2-methoxyphenyl)methyl]-2-phenyl-1-piperazinyl]ethyl]-4-phenyl-4-piperidinol.



The product from Example 8 (free form) (0.7 gram, 1.4 mmol) was used to prepare the title compound by employing a procedure analogous to the procedure for Example 4. The title compound was a solid, m.p. 63-65 °C; FAB MS $[M+1]^+$ 486; Found, C, 75.86; H, 8.52; N, 8.54. Calcd. for $C_{31}H_{39}N_3O_2$, C, 75.54; H, 8.14; N, 8.53.

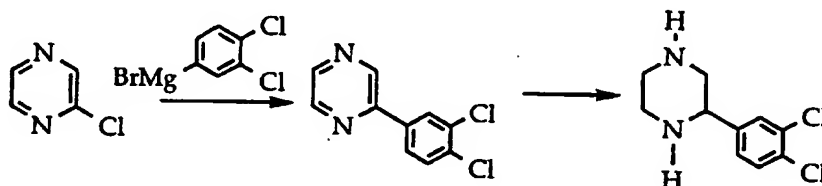
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EXAMPLE 10

2-(3,4-dichlorophenyl)piperazine

A. Synthesis:

METHOD 1



2-(3,4-Dichlorophenyl)pyrazine was prepared according to the analogous method described in Example 1. m.p. 118-119 °C; FAB MS $[M+1]^+$ 35 Cl 225.

To a solution of 2-(3,4-dichlorophenyl)pyrazine (10 g, 44.43 mmol) in dry THF (150 mL) was added slowly a solution of DIBAL-H (1M in THF, 444.3 mL) through a dropping funnel at 10 °C under N_2 . The color of solution turned into red wine at the end of addition. The solution was gradually warmed up to room temperature overnight. After completion (checked by TLC) the reaction was quenched slowly by the addition of saturated Na_2SO_4 solution until no more H_2 evolved. White precipitate

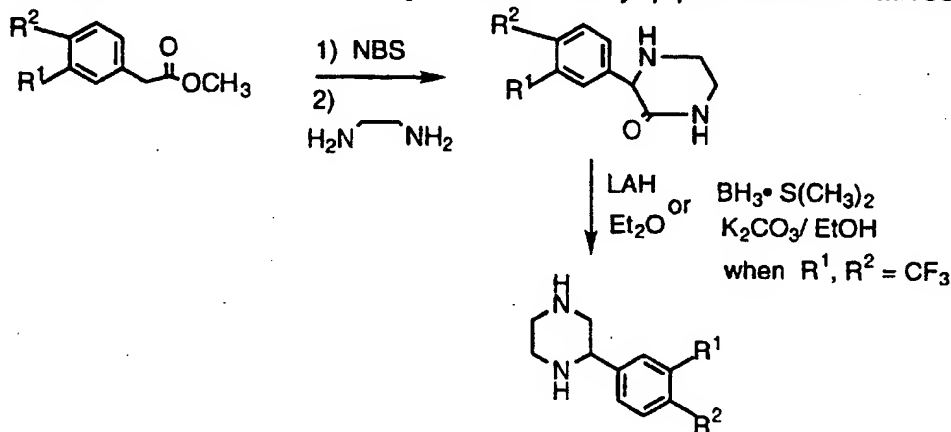
- 50 -

was formed after stirring for 1.0 h. The precipitate was filtered off, rinsed with THF, dried over MgSO_4 and evaporated to dryness. The crude material (10 g) was purified by flash chromatography on 300 g of flash grade silica gel in 7.5 % $\text{NH}_3\text{-MeOH/CH}_2\text{Cl}_2$ to give 4.11g (17.77 mmol, 40%) of 2-(3,4-dichloro-phenyl)piperazine. m.p. 74-76°C; FAB MS $[\text{M}+1]^+$ 231.

METHOD 2.

2-(3,4-Dichlorophenyl)piperazine was also synthesized according to the method published in *J. Med. Chem.*, 9, 181, 1966.

10 General method for the synthesis of 2-aryl-piperazine derivatives:



R^1 = Cl, H or other substituents i.e. OCH_3 , CF_3 , Br, I, F, etc.

R^2 = Cl, H or other substituents i.e. OCH_3 , CF_3 , Br, I, F, etc.

B. Resolution of 2-(3,4-dichlorophenyl)piperazine

Step 1

15 A solution of 2-(3,4-dichlorophenyl)piperazine (36.05 g, 0.156 mol) in methanol (200 mL) was treated with a solution containing two equivalents of N-acetyl-L-leucine (54.02 g, 0.312 mol) and heated until all of the material was dissolved. EtOAc (2.2 L) was added to this solution and allowed to stand at ambient temperature overnight. The solvent phase was decanted from the precipitated salt and concentrated in vacuo. This procedure was repeated using 37.88 g of 2-(3,4-dichlorophenyl)-piperazine (0.164 mol) and 56.68 g of N-acetyl-L-leucine (0.327 mol).

Step 2

25 The concentrated salts from both solvent phases in step 1 were combined and heated in methanol (550 mL) until all of the material dissolved. EtOAc (2.75 L) was added to this solution and allowed to stand

at ambient temperature overnight. The solvent phase was decanted from the precipitated salt and concentrated in vacuo to give ~95 g of piperazine salt (72% ee of enantiomer A).

Step 3

- 5 The salt from the solvent phase in step 2 was dissolved in a solution of H₂O (800 mL) and aq. ammonia (400 mL) and extracted with CH₂Cl₂ (4 x 400 mL). The combined organic layers were dried with MgSO₄ and concentrated to give 37 g of the piperazine free base. The free base was successively recrystallized from hexane (890, 600 and 450 mL) to give 16
10 g of piperazine (>99.9% ee of enantiomer A). $[\alpha]_D^{24.7} = -45.0^\circ$ (MEOH)

Step 4

- The precipitated salts from step 1 were combined and heated in methanol (220 mL) until all of the material dissolved. EtOAc (2.2 L) was added to this solution and allowed to stand at ambient temperature
15 overnight. The solvent phase was decanted from the precipitated salt and dried in vacuo to give ~43 g of piperazine salt (93% ee of enantiomer B).

Step 5

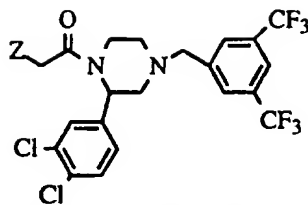
- A 12.3 g portion of salt (75% ee of enantiomer B) prepared an analogous procedure to that in step 4 was dissolved in 0.5 M NaOH (400
20 mL) and extracted with CH₂Cl₂ (4 x 155 mL). The combined organic layers were dried with MgSO₄ and concentrated to give 3.72 g of the piperazine free base. The free base was successively recrystallized from hexane (90 and 70 mL) to give 2.1 g of piperazine (98% ee of enantiomer B).

25

EXAMPLE 11

- By procedures analogous to the procedures described in Examples 1, 2 and 10 but using (+,-)-1-[[3,5-bis(trifluoromethyl)phenyl]-methyl]-3-(3,4-dichlorophenyl)-piperazine in place of (+,-)-1-[[3,5-bis(trifluoromethyl)phenyl]methyl]-3-phenyl-piperazine, and employing
30 appropriate heterocyclic reagents (Z group), listed below, the following compounds were prepared.

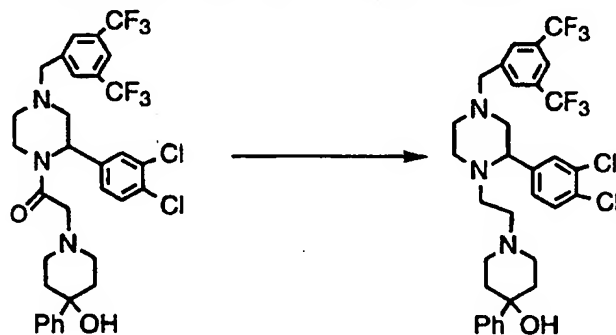
- 52 -



Z		m.p °C	[M+1] ⁺ FABMS based on ³⁵ Cl	[M+1] ⁺ High Res. MS
	free form	108-110		Cal'd 715.2041 Found 715.2050
	2 H ₂ O	58-60		Cal'd 673.1936 Found 673.1935
	free form	55-57	672	
	free form	74-76	687	
	free form	82-83	750	
	free form	105-107	702	
	free form	85-86	674	

EXAMPLE 12

(+,-)-1-[2-[4-[[3,5-bis(trifluoromethyl)phenyl]methyl]-2-(3,4-dichlorophenyl)-1-piperazinyl]ethyl]-4-phenyl-4-piperidinol.



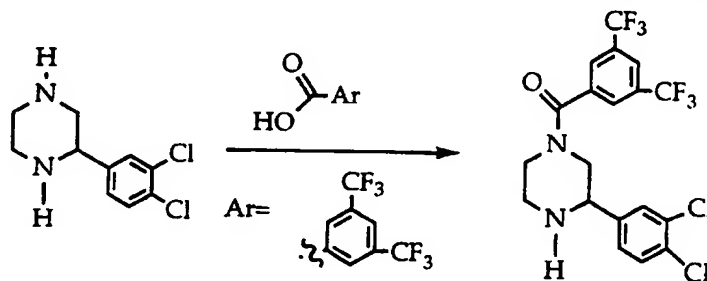
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By a procedure analogous to the procedure described in Example 4, using the last compound from Example 11, as a starting material, the title compound was prepared in 67% yield as a solid, m.p. 71-72 °C; FAB MS [M+1]⁺ 660; Found, C, 58.08; H, 5.14; N, 6.40; F, 17.37. Calcd. for C₃₂H₃₃N₃Cl₂F₆O, C, 58.19; H, 5.04; N, 6.36; F, 17.26.

10

EXAMPLE 13

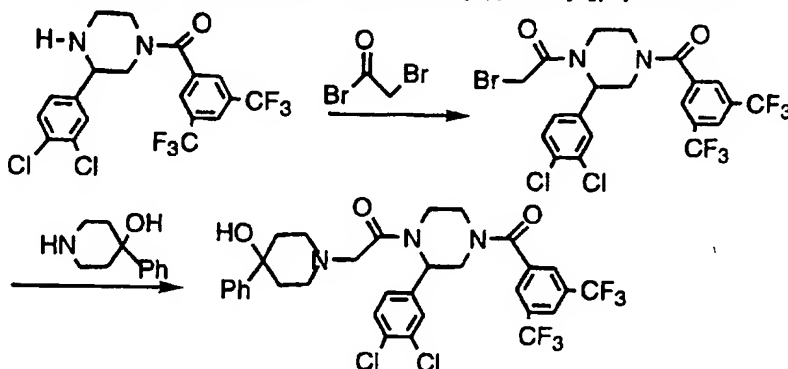
(+,-)-[3,5-bis(trifluoromethyl)benzoyl]-3-(3,4-dichlorophenyl)piperazine



To a cooled solution of CH_2Cl_2 (103 mL) containing 2-(3,4-dichlorophenyl)piperazine (1.15 g 5.0 mmol), 3,5-bis-(trifluoromethyl)-benzoic acid (1.34 g, 5.09 mmol), and N-hydroxybenzotriazole monohydrate (0.688 g, 5.09 mmol) at -20°C were added Et_3N (0.711 mL 5.09 mmol) and N,N-dimethylaminopropyl ethylcarbodiimide (DEC) (0.967 g, 5.09 mmol) under nitrogen. The reaction was kept at -20°C for an hour and gradually warmed to RT overnight. After stirring 20 h, the reaction was complete and CH_2Cl_2 (200 mL) was added. The organic solution was washed with 5% NaHCO_3 (80 mL) and brine (80 mL, 2x), dried over MgSO_4 , filtered and concentrated under vacuum to give 2.1 g of crude product. The product was purified by flash chromatography on flash grade silica gel (120 g), eluting with 2% NH_3 -MeOH/ CH_2Cl_2 to give a foam solid (1.25 g, 2.65 mmol, 53%). m.p. 50 - 53°C ; FAB MS $[\text{M}+1]^+$ 470.9; Calcd. for $\text{C}_{19}\text{H}_{14}\text{ON}_2\text{F}_6\text{Cl}_2$, C, 48.42; H, 2.99; N, 5.94; F, 24.19; Cl, 15.05. Found, C, 48.57; H, 2.90; N, 5.94; F, 23.90; Cl, 15.03.

EXAMPLE 14

(+,-)-4-[3,5-bis(trifluoromethyl)benzoyl]-2-(3,4-dichlorophenyl)-1-[[4-hydroxy-4-phenyl-1-piperidiny]]acetyl]piperazine



To a solution of (+,-)-[3,5-bis(trifluoromethyl)benzoyl]-3-(3,4-dichlorophenyl)piperazine (0.6 g, 1.274 mmol) in dry CH_2Cl_2 (12.0 mL) at

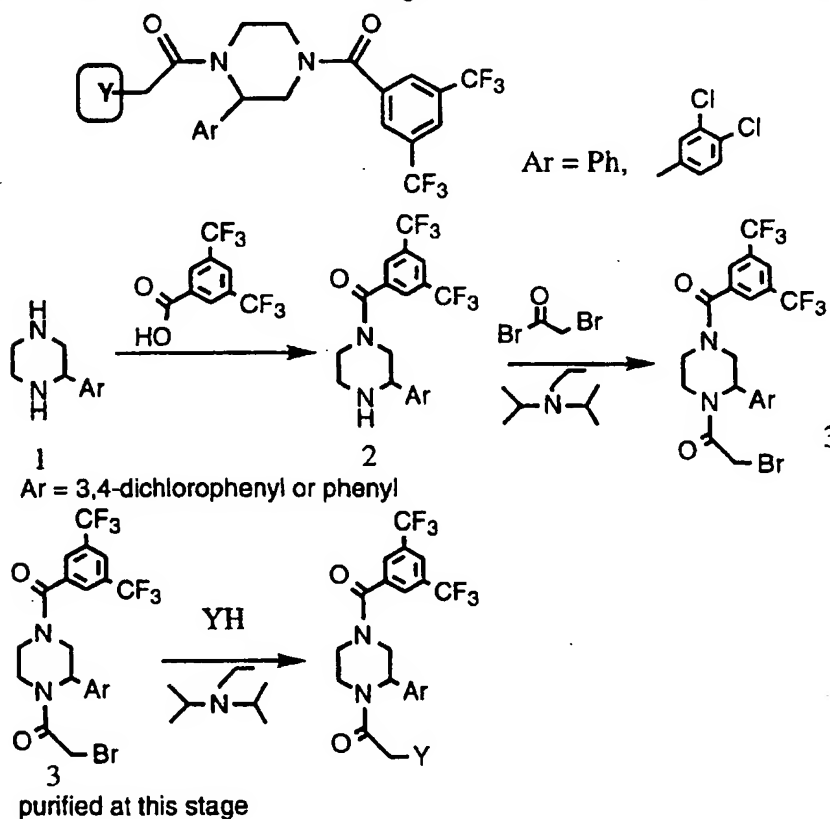
- 54 -

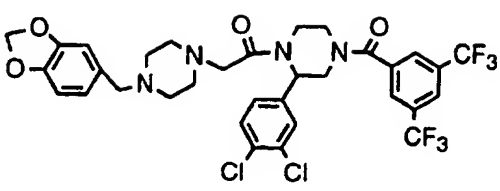
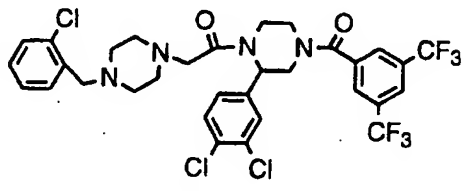
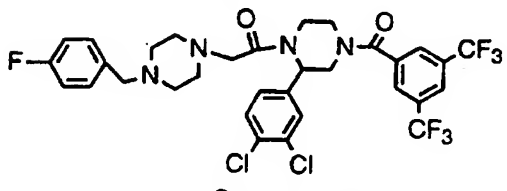
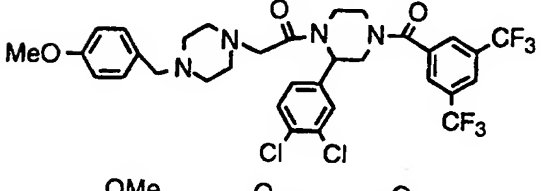
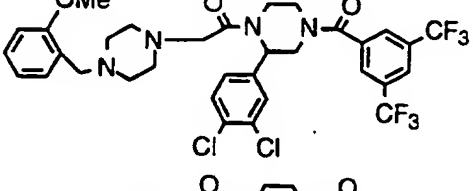
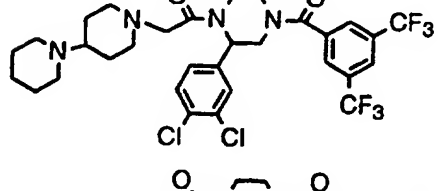
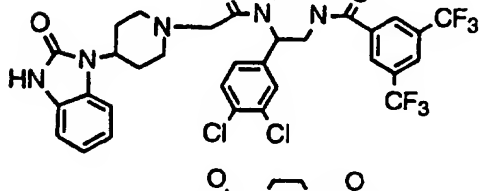
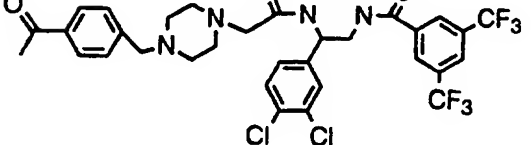
-78 °C was added diisopropylethylamine (0.266 mL, 1.53 mmol) followed by the dropwise addition of bromoacetyl bromide (0.124 mL, 1.40 mmol). After stirring at -78 °C for 3.5 h under nitrogen, additional diisopropyl ethylamine (0.234 mL, 1.342 mmol) and 4-amino-1-benzyl piperidine (0.279 mL, 1.342 mmol) were added at -78 °C. The reaction was gradually warmed to RT overnight. After the reaction was complete, the reaction was diluted with CH₂Cl₂ (200 mL), washed with brine (80 mL, 3x) and dried over MgSO₄. After filtration, the solvent was removed under vacuum to give a light yellow solid which was purified by flash chromatography on flash grade silica gel (150 g), eluting with 5% NH₃-MeOH/CH₂Cl₂ to give the title compound was prepared in 72 % yield as a solid, m.p. 104-106°C, FAB MS [M+1]⁺ 688.1

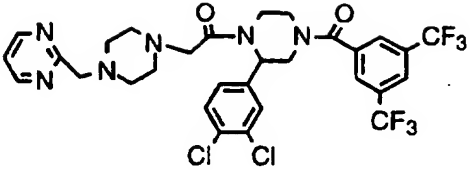
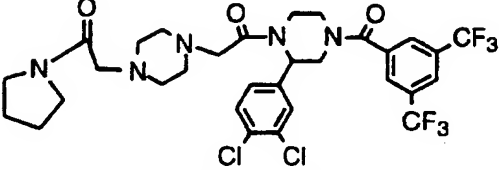
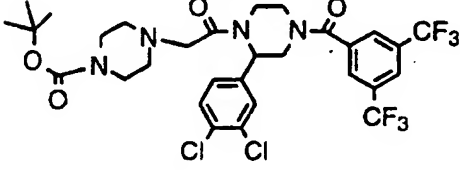
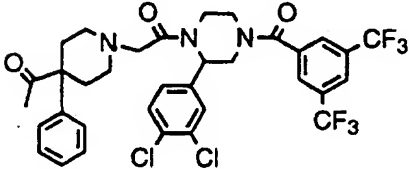
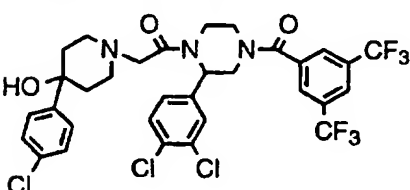
Calcd. for C₃₂H₂₉N₃O₃F₆Cl₂ • 0.25 H₂O, C, 55.45; H, 4.30; N, 6.06; F, 16.45; Cl, 10.23. Found, C, 55.40; H, 4.38; N, 6.05; F, 16.83; Cl, 10.63.

EXAMPLE 15

By employing methods analogous to those described in Example 13 and Example 14 using appropriate alkylation reagents, the following compounds were obtained according to the schemes shown below.



	FAB MS [M+1] ⁺
	731.0
	707.0
	691.0
	703.0
	703.0
	679.0
	728.0
	715.1

	FAB MS [M+1] ⁺
	675.0
	708.0
	697.1
	714.0
	724.0

EXAMPLE 16

(+,-)-2-(3,4-dichlorophenyl)-4-(2-methoxybenzoyl)-1-[[4-(4-hydroxy-4-phenyl-1-piperidiny)]acetyl]piperazine

- 5 By employing methods analogous to those described in Example 13 and Example 15 and using 2-methoxybenzoic acid in place of bis(3,5-trifluoromethyl) benzoic acid the title compound was prepared in 71% yield as a solid, m.p. 112-114°C, FAB MS [M+1]⁺ 582.0.

EXAMPLE 17

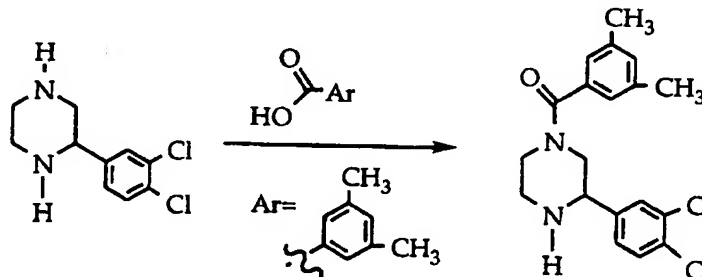
- 10 (+,-)-4-[3,5-bis(trifluoromethyl)benzoyl]-2-phenyl-1-[[4-(4-hydroxy-4-phenyl-1-piperidiny)]acetyl]piperazine

- By employing methods analogous to those described in Example 13 and Example 15, and using 2-phenylpiperazine in place of 2-(3,4-dichlorophenyl)-piperazine the title compound was prepared in 90 %
 15 yield as a solid: m.p. 101-102 °C, FAB MS [M+1]⁺ 620.4;

Calcd. for $C_{32}H_{29}N_3O_3F_6Cl_2 \cdot 0.25 H_2O$, C, 61.57; H, 5.09; N, 6.73; F, 18.27. Found, C, 61.41; H, 5.08; N, 6.71; F, 18.28.

EXAMPLE 18

(+,-)-[3,5-dimethylbenzoyl]-3-(3,4-dichlorophenyl)piperazine



5

To a cooled solution of CH_2Cl_2 (600 mL) containing 2-(3,4-dichlorophenyl)piperazine (6.934 g, 30 mmol), 3,5-dimethylbenzoic acid (4.55 g, 30 mmol), and N-hydroxybenzotriazole monohydrate (4.05 g, 30 mmol) at $-20^\circ C$ were added Et_3N (4.2 mL, 30 mmol) and N,N-dimethylaminopropylethylcarbodiimide (DEC) (5.86 g, 30 mmol) under nitrogen. The reaction was kept at $-20^\circ C$ for an hour and gradually warmed to RT overnight. After stirring 22 h, the reaction was complete and CH_2Cl_2 (200 mL) was added. The organic solution was washed with brine (150 mL, 3x), dried over $MgSO_4$, filtered and concentrated under vacuum to give 8.2 g of crude product. The product was crystallized from CH_2Cl_2 /Hexane to give a light yellow solid (6.3 g, 17.34 mmol, 57.8%), m.p. $139-141^\circ C$; FAB MS $[M+1]^+$ 363.1.

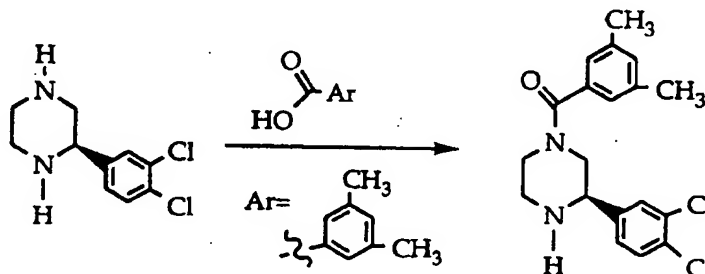
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EXAMPLE 19

(+)-[3,5-dimethylbenzoyl]-3-(R)-(3,4-dichlorophenyl)piperazine
(Enantiomer B)

20



The title compound was prepared by an analogous method to that described in Example 18 using (-)-2-(R)-(3,4-dichlorophenyl)piperazine in place of (+,-)-2-(3,4-dichlorophenyl)piperazine, m.p. $97-100^\circ C$; FAB MS $[M+1]^+=363.1$;

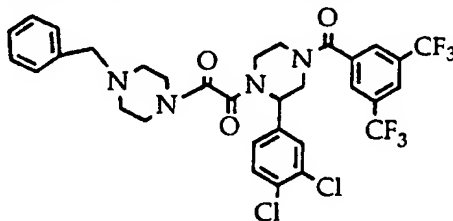
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$[\alpha]_D^{22.5} = +87.2^\circ$ (MeOH).

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EXAMPLE 20

(+,-)-4-[3,5-bis(trifluoromethyl)benzoyl]-2-(3,4-dichlorophenyl)-
1[1,2-dioxo-2-[4-(phenylmethyl)-1-piperaziny]ethyl]piperazine



5 Step 1 A solution of 1-benzylpiperazine (1.75 mL, 10 mmol) in CH_2Cl_2 (60 mL) was treated with methyl acrylate (0.90 mL, 10 mmol) at ambient temperature for 2 days. The reaction mixture was concentrated to give a clean Michael product (2.6 g, 10 mmol, 100%).

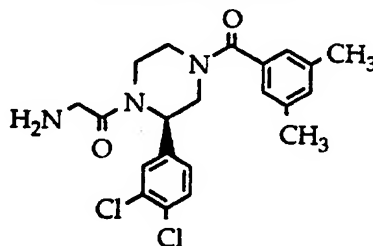
Step 2 A solution of the product from step 1 (0.92 g, 3.5 mmol) in
10 methanol (10 mL) was treated with 1 M LiOH (5.2 mL, 5.2 mmol) for 1 h. The reaction mixture was concentrated in vacuo at 50°C and the residue was suspended in CH_2Cl_2 (25 mL), filtered and concentrated to obtain 0.72 g of the desired acid (2.9 mmol, 83%).

Step 3 A solution of the product from step 2 (0.72 g, 2.9 mmol) in
15 benzene (12 mL) was treated with oxalyl chloride (300 mL, 3.4 mmol) and DMF (1 drop) at 0°C . The reaction mixture was warmed up to ambient temperature and stirred for 90 min. The reaction mixture was concentrated in vacuo and the residue was suspended in CH_2Cl_2 (20 mL). To this suspension were added 1-[3,5-bis(trifluoro-methyl)benzoyl]-
20 3-(3,4-dichlorophenyl)piperazine (1.4 g, 2.9 mmol) in CH_2Cl_2 (14 mL) and triethylamine (0.80 mL, 5.7 mmol). The reaction mixture was stirred for 1 h and H_2O (100 mL) was added. The mixture was extracted with CH_2Cl_2 (50 and 25 mL). The combined organic layers were washed with brine (50 mL), dried (MgSO_4) and concentrated. The crude material was
25 purified by silica gel chromatography, eluanting with CH_2Cl_2 :methanol (50:1) to obtain 0.64 g (0.91 mmol, 31%) of the titled compound as a colorless foam.

HRMS (FAB, $\text{M}+\text{H}^+$): m/e calc'd for $[\text{C}_{32}\text{H}_{29}\text{Cl}_2\text{F}_6\text{N}_4\text{O}_3]^+$: 701.1521, found 701.1513.

30

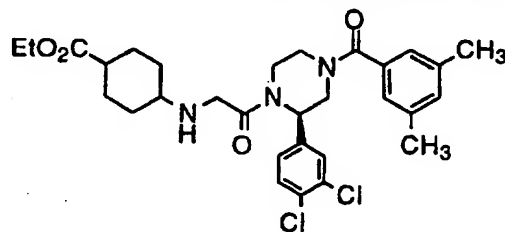
EXAMPLE 21

1-[3,5-dimethylbenzoyl]-3-(R)-(3,4-dichlorophenyl)piperazine
(enantiomer B)

- 5 To a solution of BOC glycine (0.918 g, 5.24 mmol), (+)-(3,5-dimethylbenzoyl)-3-(R)-(3,4-dichlorophenyl)piperazine (enantiomer B) (1.80 g, 4.95 mmol) (prepared in Example 19), DEC (0.983 g, 5.13 mmol), HOBT (0.687 g, 5.07 mmol) and Hünig's base (0.92 mL, 5.1 mmol) in CH₂Cl₂ (100 mL) was left stirring for 2.5 d. The reaction mixture was added to CH₂Cl₂ (200 mL) and washed with sat. NaHCO₃ (3 x 100 mL), brine (100 mL), dried with MgSO₄ and concentrated. The crude material was treated with MeOH saturated HCl (25 mL) for 10 h and concentrated. The resulting residue suspended in 0.3 N NaOH (150 mL) and extracted with CH₂Cl₂, (3 x 50 mL). The combined organic layers were washed with brine (50 mL), dried with MgSO₄, concentrated, and purified by flash chromatography on silica gel, eluting with 20:1:0.2 CH₂Cl₂/MeOH/conc. aq. NH₃ to give 0.95 g of the
- 10
- 15
- 20
- 25
- titled product as a white solid. HRMS (FAB, M+H⁺); m/e calc'd [C₃₁H₂₄Cl₂N₃O₂]⁺: 420.1246 found 420.1254.

EXAMPLE 22

2-(R)-(3,4-dichlorophenyl)-4-(3,5-dimethylbenzoyl)-1-[[4-carbethoxycyclohexyl)amino]acetyl]piperazine (enantiomer B)



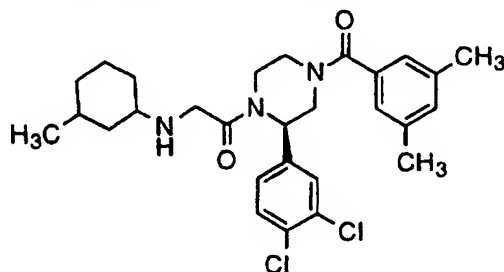
- A CH₂Cl₂ (2.0 mL) solution containing the amino compound from Example 21 (0.10 g, 0.23 mmol) and 1-carbethoxy-4-piperidone (39 mg, 0.23 mmol) was treated with NaBH(OAc)₃ (63 mg, 0.32 mmol) and acetic acid (15 mL, 0.26 mmol) and left stirring overnight. The reaction mixture was quenched with 1 N NaOH and extracted with CH₂Cl₂ (50 mL, 3 x). The combined organic layers were washed with brine, dried with

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MgSO₄ and chromatographed on silica gel eluting with 5% NH₃ sat. MeOH/CH₂Cl₂ to give 46 mg of the titled product as a white solid. HRMS (FAB, M+H⁺); m/e calc'd [C₃₀H₃₈Cl₂N₃O₄]⁺: 574.2239, found 574.2250.

EXAMPLE 23

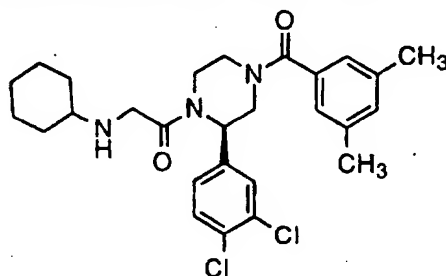
- 5 2-(R)-(3,4-dichlorophenyl)-4-(3,5-dimethylbenzoyl)-1-[[[(3-methylcyclohexyl)amino)acetyl]piperazine (diastereomers from enantiomer B)



- By employing method analogous to that described in Example 22 using 3-methylcyclohexanone, the title compound was obtained. HRMS (FAB, M+H⁺); m/e calc'd [C₂₈H₃₆Cl₂N₃O₂]⁺: 516.2185, found 516.2199.

EXAMPLE 24

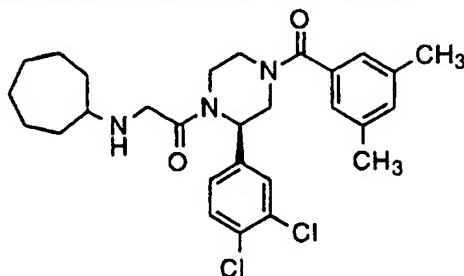
- 1-[(cyclohexylamino)acetyl]-2-(R)-(3,4-dichlorophenyl)-4-(3,5-dimethylbenzoyl) piperazine (enantiomer B)



- 15 By employing method analogous to that described in Example 22 using cyclohexanone, the title compound was obtained. HRMS (FAB, M+H⁺); m/e calc'd [C₂₇H₃₄Cl₂N₃O₂]⁺: 502.2028, found 502.2025.

EXAMPLE 25

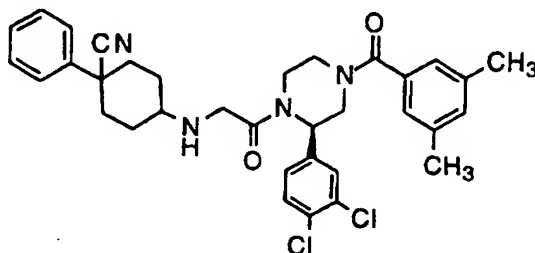
- 20 1-[(cycloheptylamino)acetyl]-2-(R)-(3,4-dichlorophenyl)-4-(3,5-dimethylbenzoyl) piperazine (enantiomer B)



By employing method analogous to that described in Example 22 using cycloheptanone, the title compound was obtained. HRMS (FAB, $M+H^+$); m/e calc'd $[C_{28}H_{36}Cl_2N_3O_2]^+$: 516.2185, found 516.2177.

EXAMPLE 26

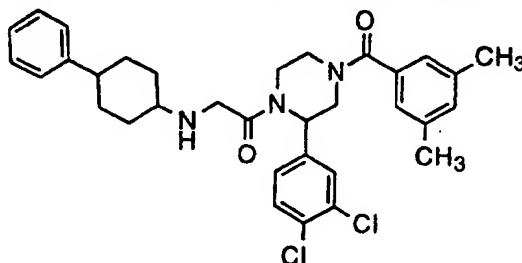
- 5 1-[[[(4-cyano-4-phenylcyclohexyl)amino]acetyl]-2-(R)-(3,4-dichlorophenyl)-4-(3,5-dimethylbenzoyl)piperazine (enantiomer B)



- By employing method analogous to that described in Example 22 using 4-cyano-4-phenylcyclohexanone, the title compound was obtained.
 10 HRMS (FAB, $M+H^+$); m/e calc'd $[C_{34}H_{37}Cl_2N_4O_2]^+$: 603.2294, found 603.2271.

EXAMPLE 26A

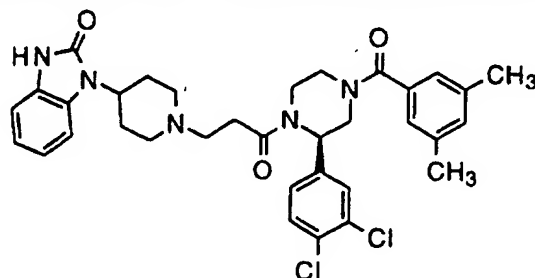
- (+/-)-2-(3,4-dichlorophenyl)-4-(3,5-dimethylbenzoyl)-1-[[[(4-phenylcyclohexyl) amino]acetyl]-piperazine



- 15 By employing methods analogous to that described in Examples 21 and 22 using the (+,-)-(3,5-dimethylbenzoyl)-3-(3,4-dichlorophenyl)-piperazine (enantiomer B) from example 18 and 4-phenylcyclohexanone, the title compound was obtained. HRMS (FAB, $M+H^+$); m/e calc'd
 20 $[C_{33}H_{39}Cl_2N_3O_2]^+$: 578.2341, found 578.2327.

EXAMPLE 27

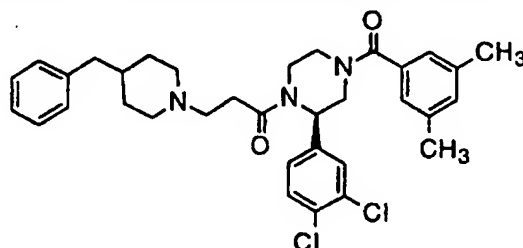
2-(R)-(3,4-dichlorophenyl)-4-(3,5-dimethylbenzoyl)-1-[3-[4-(2-keto-1-benzimidazolyl)piperidinyl]-1-oxopropyl]piperazine (enantiomer B)



- 5 A cooled CH_2Cl_2 (4 mL) solution containing (+)-(3,5-dimethylbenzoyl)-3-(3,4-dichlorophenyl)piperazine (enantiomer B) from Example 19 (153 mg, 0.42 mmol) and Hünig's base (0.12 mL, 0.86 mmol) at -78°C was treated with 3-chloropropionyl chloride (0.040 mL, 0.42 mmol) and warmed to RT for 1 h. The reaction mixture was then concentrated in
- 10 vacuo and resuspended in acetonitrile (2 mL). To the reaction mixture was added 4-(2-keto-1-benzimidazolyl)piperidine (104 mg, 0.48 mmol) and left heating overnight at 70°C . The reaction mixture was concentrated and purified by silica gel chromatography, eluting with 20:1:0.1 $\text{CH}_2\text{Cl}_2/\text{MeOH}/\text{conc. aq. NH}_3$ to give 104 mg of the title
- 15 compound as a white solid (39 % yield). HRMS (FAB, $\text{M}+\text{H}^+$); m/e calc'd $[\text{C}_{34}\text{H}_{38}\text{Cl}_2\text{N}_5\text{O}_3]^+$: 634.2352, found 634.2351.

EXAMPLE 28

2-(R)-(3,4-dichlorophenyl)-4-(3,5-dimethylbenzoyl)-1-[3-[4-(phenylmethyl)-1-piperidinyl]-1-oxopropyl]piperazine (enantiomer B)



20

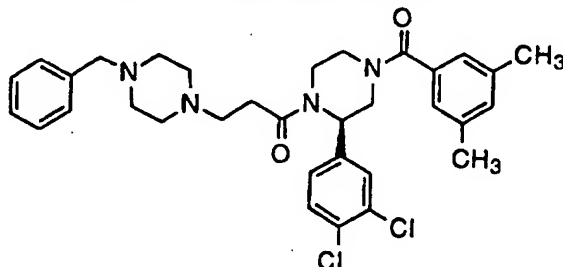
By employing method analogous to that described in Example 27 using 4-benzylpiperidine, the title compound was obtained. HRMS (FAB, $\text{M}+\text{H}^+$); m/e calc'd $[\text{C}_{34}\text{H}_{40}\text{Cl}_2\text{N}_3\text{O}_2]^+$: 592.2498, found 592.2494.

25

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EXAMPLE 29

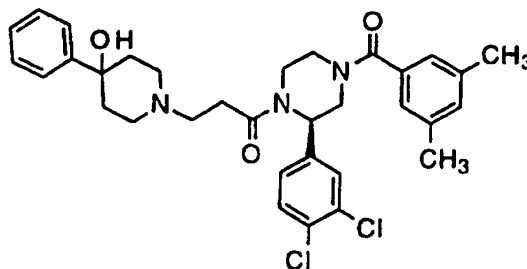
2-(R)-(3,4-dichlorophenyl)-4-(3,5-dimethylbenzoyl)-1-[1-oxo-3-[4-(phenylmethyl)-1-piperidiny]propyl]piperazine (enantiomer B)



5 By employing method analogous to that described in Example 27 using 1-benzyl-piperazine, the title compound was obtained. HRMS (FAB, M+H⁺); m/e calc'd [C₃₃H₄₉Cl₂N₄O₂]⁺: 593.2450, found 593.2464.

EXAMPLE 30

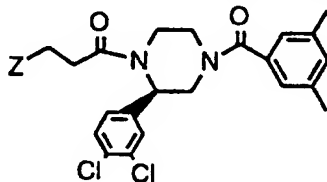
10 2-(R)-(3,4-dichlorophenyl)-4-(3,5-dimethylbenzoyl)-1-[3-[4-hydroxy-4-phenyl-1-piperidiny]-1-oxopropyl]piperazine (enantiomer B)



By employing method analogous to that described in Example 27 using 4-hydroxy-4-phenylpiperidine, the title compound was obtained. HRMS (FAB, M+H⁺); m/e calc'd [C₃₃H₃₈Cl₂N₃O₃]⁺: 594.2290, found 594.2285.

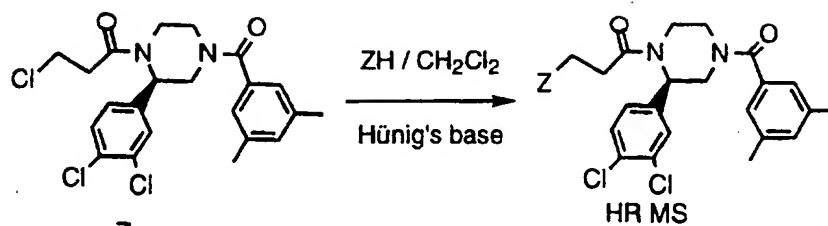
EXAMPLE 31

The following examples were prepared analogously to the methods described in Example 27 by using appropriate piperidine or piperazine derivative in place of 4-(2-keto-1-benzimidazolyl)piperidine.



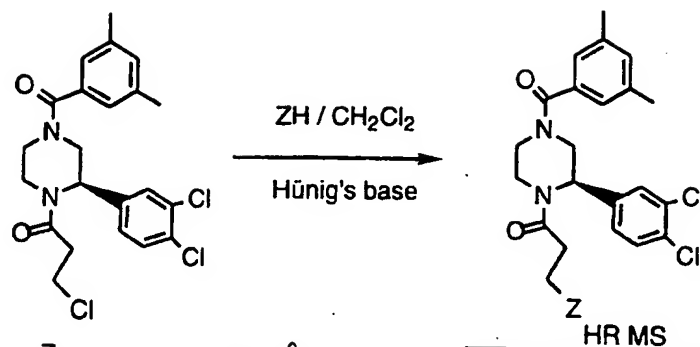
20

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Z	measured	calculated
	593.2464	593.2450
	592.2494	592.2498
	594.2285	594.2290
	634.2351	634.2352
	575.2183	575.2192
	607.2612	607.2607
	579.2284	579.2294
	580.2251	580.2246
	609.2402	609.2399
	608.2450	608.2447
	599.2550	599.2556

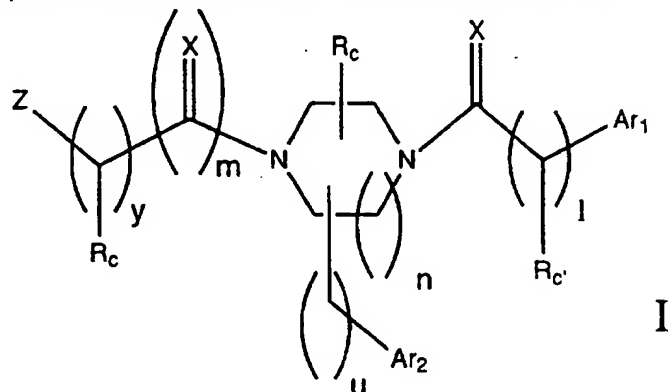
- 65 -



Z	m.p. °C	HR MS	
		measured	calculated
	112-117	585.2765	585.2763
	223-226	643.3179	643.3182
	187 decomposed	629.2486	629.2484
	127-130	599.2545	599.2556
	159-163	684.3070	684.3083
		578.2325	578.2341
	97-99	698.3232	698.3240
	94-97	670.3290	670.3291
	78-80	670.2924	670.2927
	82-84	581.2183	581.2199

WHAT IS CLAIMED IS:

1. A compound represented by the structural formula



5

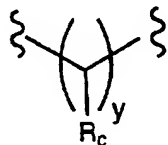
each X is independently, O, (H,H), NR_d, or S;

n is 0 to 2; u is 0 to 2; l is 0 to 2;

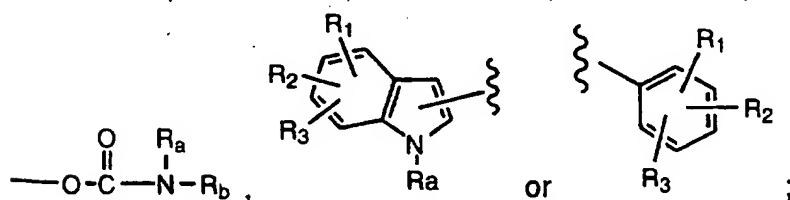
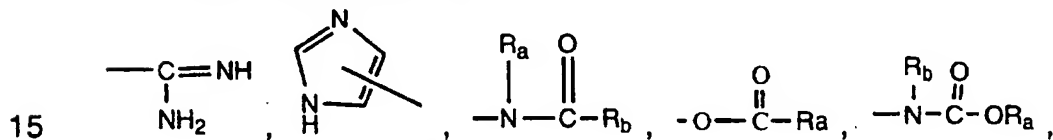
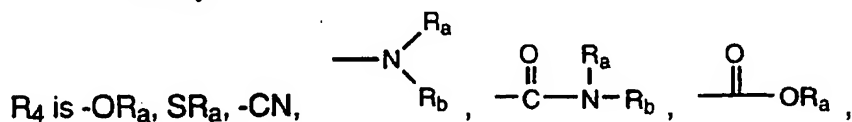
m is 1, and y is 1 to 3; or m is 2, and y is 0;

each R_c is independently H, C₁-C₆ alkyl, -(CH₂)_{n1}-R₄ where n₁ is 1 to 6;

- 10 R_d is independently selected from the group consisting of H, C₁-C₆ alkyl, CN, OR_a, phenyl, substituted phenyl, benzyl, substituted benzyl, or allyl, and with the further proviso that no more than one R_c is other than H in the



moiety;



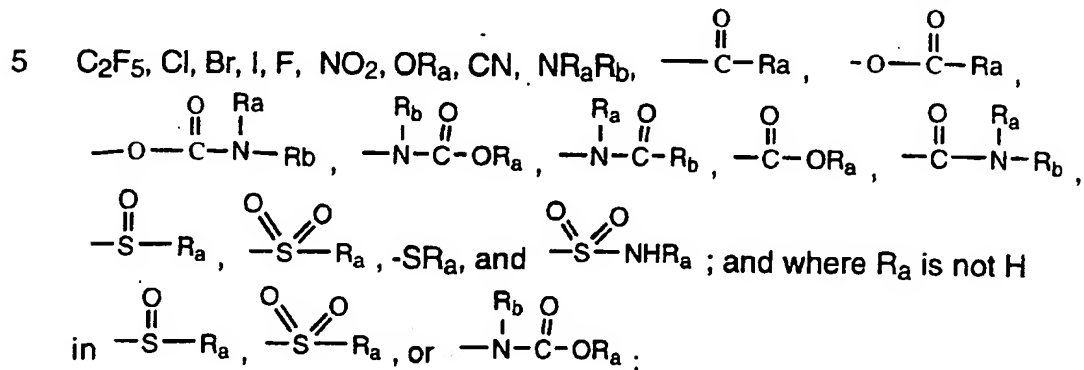
R_c' is H, C₁-C₆ alkyl or (CH₂)_nOR_a, with the proviso that no more than one R_c' is other than H;

- 20 each R_a and R_b is independently selected from the group consisting of H, C₁-C₆ alkyl, phenyl, substituted phenyl, benzyl, substituted benzyl,

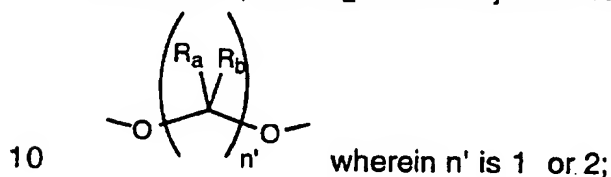
- 67 -

allyl; or when R_a and R_b are attached to the same nitrogen, then R_a and R_b together with the nitrogen to which they are attached, form a 4 to 7 member ring;

wherein each R_1 and R_2 is independently H, C_1 - C_6 alkyl, CF_3 ,

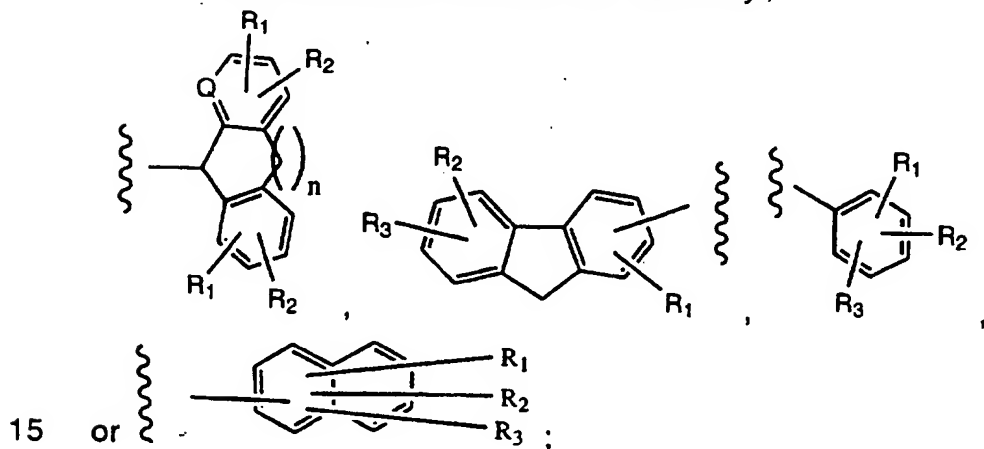


or when R_1 and R_2 are on adjacent carbons on a ring, they can form



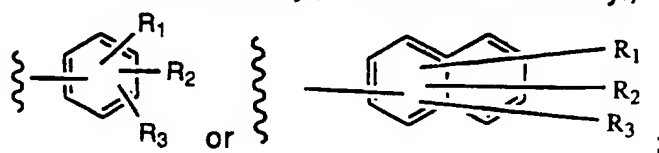
and each R_3 is independently H, C_1 - C_6 alkyl, CF_3 , C_2F_5 , $\text{---}\overset{\text{O}}{\parallel}\text{C---}Ra$,
 $\text{---}O\text{---}\overset{\text{O}}{\parallel}\text{C---}Ra$, $\text{---}\overset{\text{O}}{\parallel}\text{C}\overset{Ra}{\underset{|}{N}}\text{---}Rb$, Cl, Br, I, F, OR_a , OCF_3 , or phenyl;

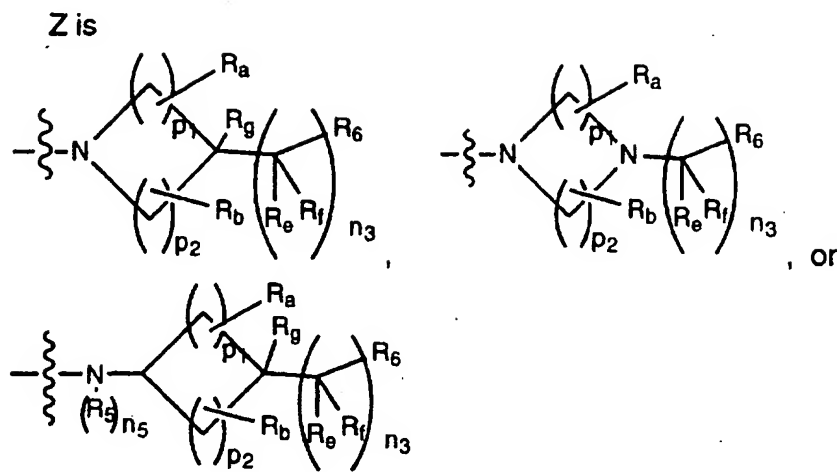
Ar_1 is heteroaryl or substituted heteroaryl,



Q is N or CH;

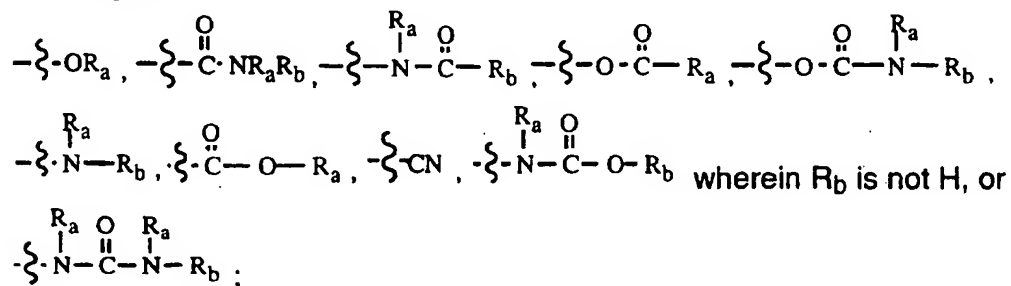
Ar_2 is heteroaryl, substituted heteroaryl,





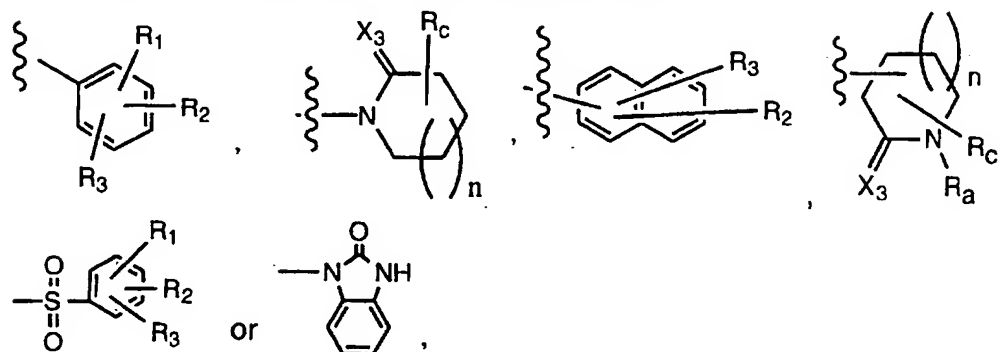
- wherein p_1 and p_2 are each independently 1 to 4 with the proviso that p_1 and p_2 added together are 2 to 6; n_5 is 1 to 2; each R_5 is independently selected from the group consisting of H, OH, $\text{—}\overset{\text{O}}{\parallel}\text{C—R}_a$, $\text{C}_1\text{—C}_6$ alkyl, $\text{—}(\text{CH}_2)_{n_1}\text{—R}_4$ where n_1 is 1 to 6 with the proviso that when n_1 is 1, R_4 is not OH or NR_aR_b ; also with the proviso that when n_5 is 2, R_5 is $\text{C}_1\text{—C}_6$ alkyl, and two R_5 can be attached to the nitrogen to form a quaternary salt;
- each R_a and R_b is independently selected from the group consisting of H, $\text{C}_1\text{—C}_6$ alkyl, phenyl, substituted phenyl, benzyl, substituted benzyl, allyl;
- n_3 is 0-4;
- each R_e and R_f is independently selected from the group consisting of H, $\text{C}_1\text{—C}_6$ alkyl, phenyl, substituted phenyl, benzyl, substituted benzyl, allyl; or R_e and R_f taken together with the carbon to which they are attached can also form a carbonyl group with the proviso that no more

- than one carbonyl group is in the moiety;
- R_g is H,

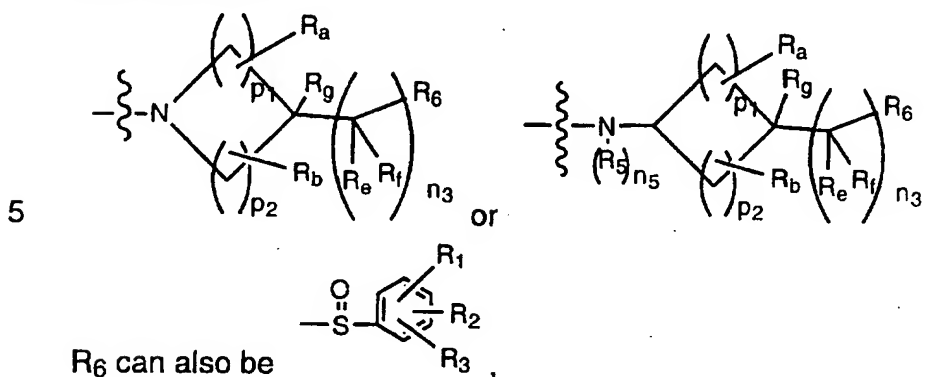


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R_6 is H, C_1 - C_6 alkyl, allyl, C_3 - C_6 cycloalkyl,



and when Z is



R_6 can also be

wherein X_3 is O, (H,H), NR_d , or S; or R_6 is heteroaryl, substituted heteroaryl, heterocycloalkyl, substituted heterocycloalkyl, when n_3 is 0-4; or when R_e , R_f taken together with the carbon atom to which they are attached form a carbonyl group and n_3 is 1, R_6 can also be OR_a wherein R_a is not H, and R_6 can also be $-(NR_a, R_b)$, O-heteroaryl, O-substituted heteroaryl, O-heterocycloalkyl, O-substituted heterocycloalkyl, $-NR_a$ -heteroaryl, $-NR_a$ -substituted heteroaryl, $-NR_a$ -heterocycloalkyl, $-NR_a$ -substituted heterocycloalkyl.

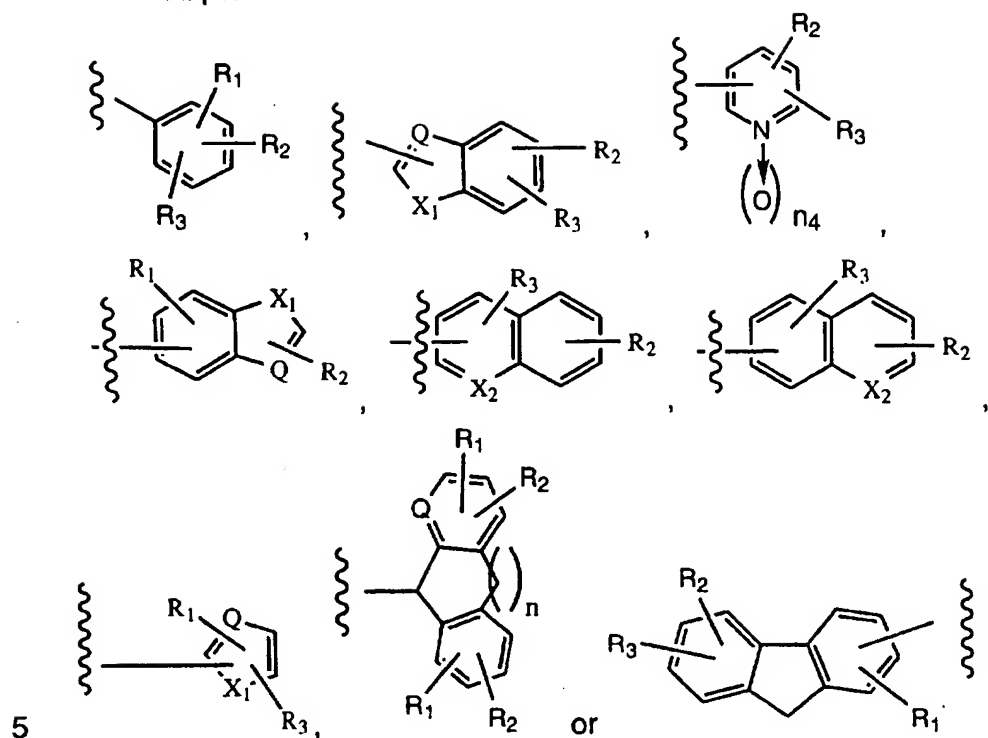
or a pharmaceutically acceptable salt thereof.

2. A compound according to claim 1, wherein each X is O or (H,H) and at least one X is O.

3. A compound according to claim 2, wherein both X's are O.

4. A compound according to any of claims 1, 2 or 3 wherein l is 0; m is 1; y is 1 to 3; n is 1; u is 0;

Ar₁ is



wherein Q is N or CH;

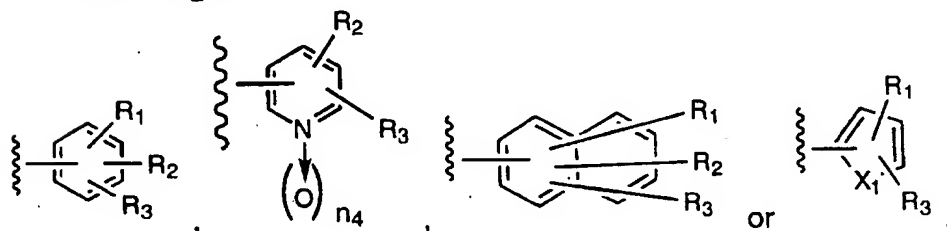
each X_1 is independently O, S or NR_a ;

each X_2 is independently CH or N; and

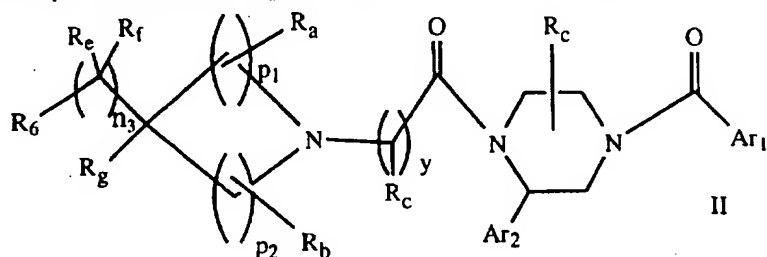
n_4 is 0 or 1;

10

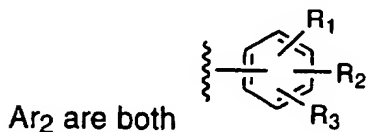
and Ar_2 is



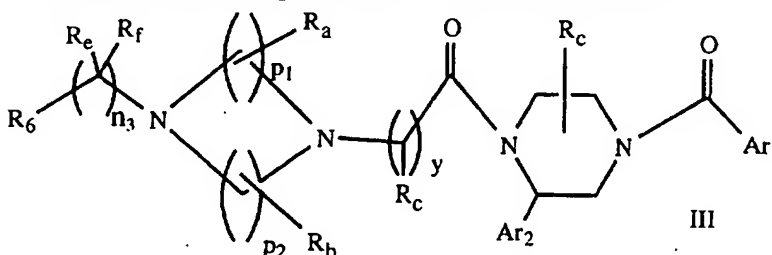
5. A compound according to claim 1 of the formula II



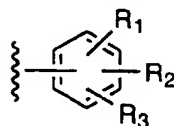
wherein R_c is H; y is 1-3; p_1 and p_2 are 2; R_e and R_f are H, C₁-C₆ alkyl, C₃-C₆ cycloalkyl or allyl; n_3 is 0-4; R_g is defined in claim 1; and Ar_1 and



- 5 6. A compound according to claim 1 of the formula III

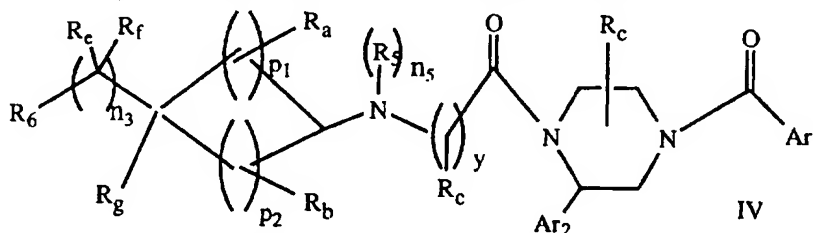


wherein R_c is H; y is 1-3; p_1 and p_2 are 2; n_3 is 0-4; R_e and R_f are H, C₁-C₆ alkyl, C₃-C₆ cycloalkyl or allyl; and Ar_1 and Ar_2 are both

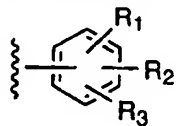


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7. A compound according to claim 1 of the formula IV

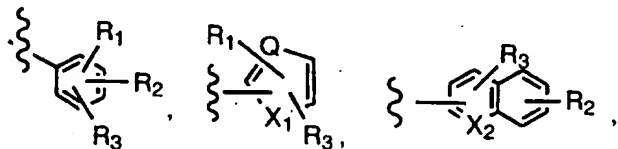


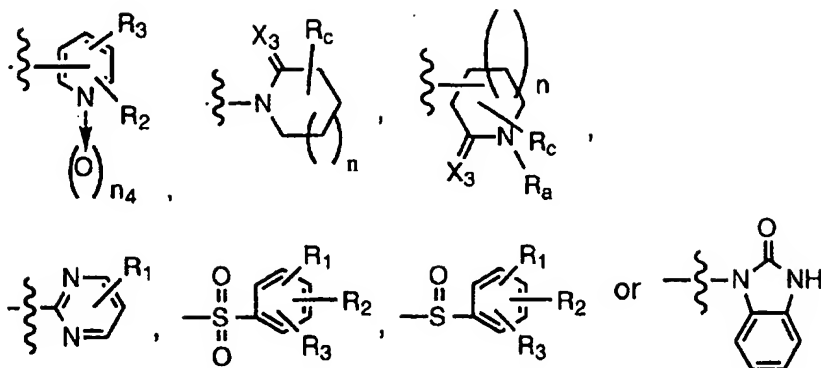
where in R_c is H; y is 1-3; p_1 , p_2 are 1-2; n_3 is 0 to 4; R_e and R_f are H, C₁-C₆ alkyl, C₃-C₆ cycloalkyl or allyl; and Ar_1 and Ar_2 are both



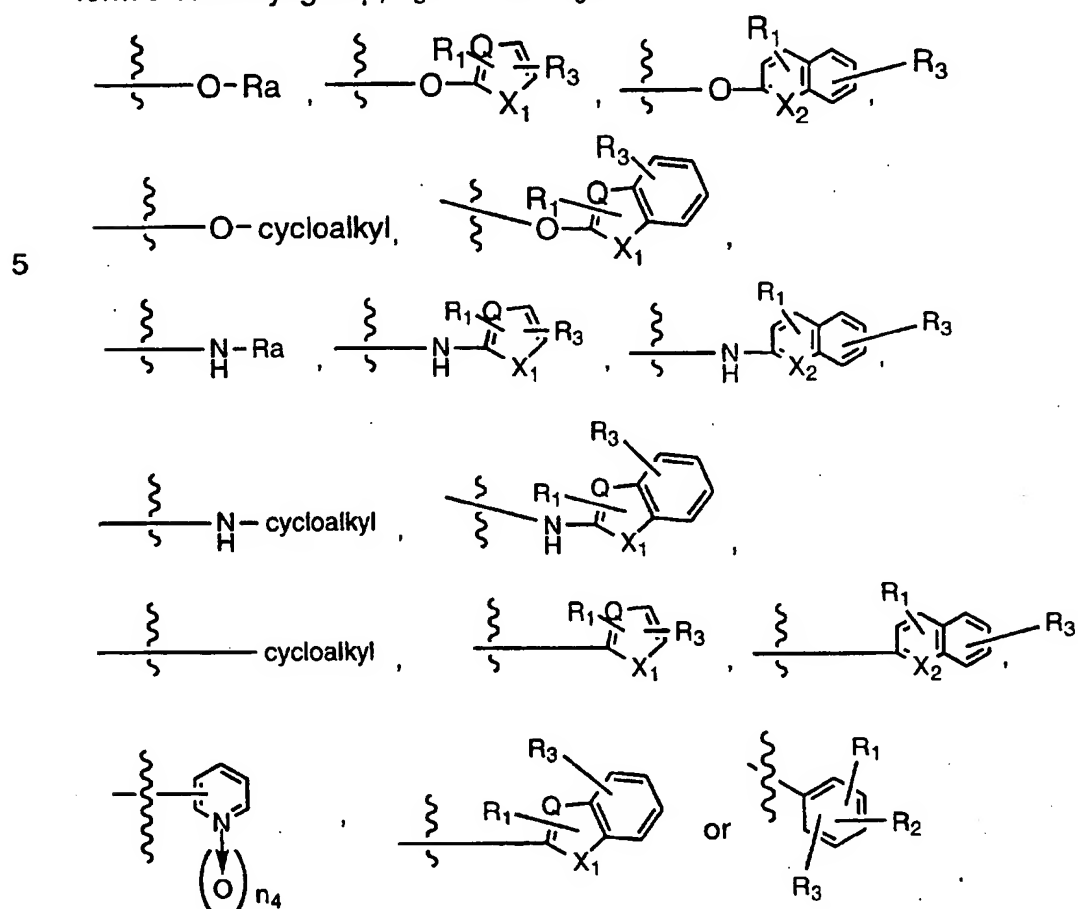
15

8. A compound according to claim 5 or 7 wherein R_6 is

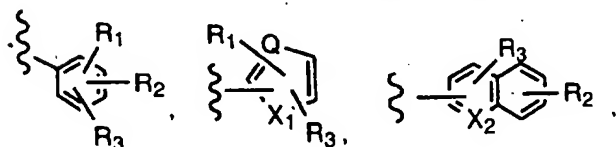




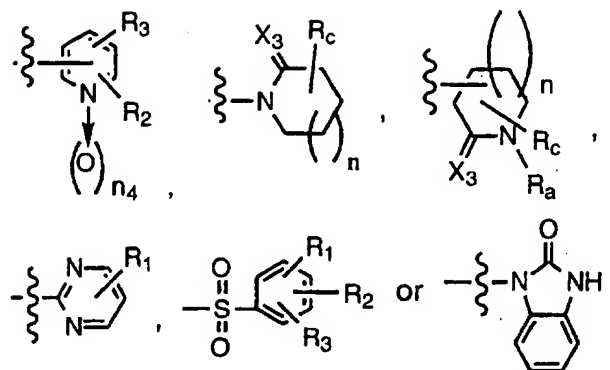
or R_E and R_F taken together with the carbon to which they are attached form a carbonyl group, n_3 is 1 and R_6 is



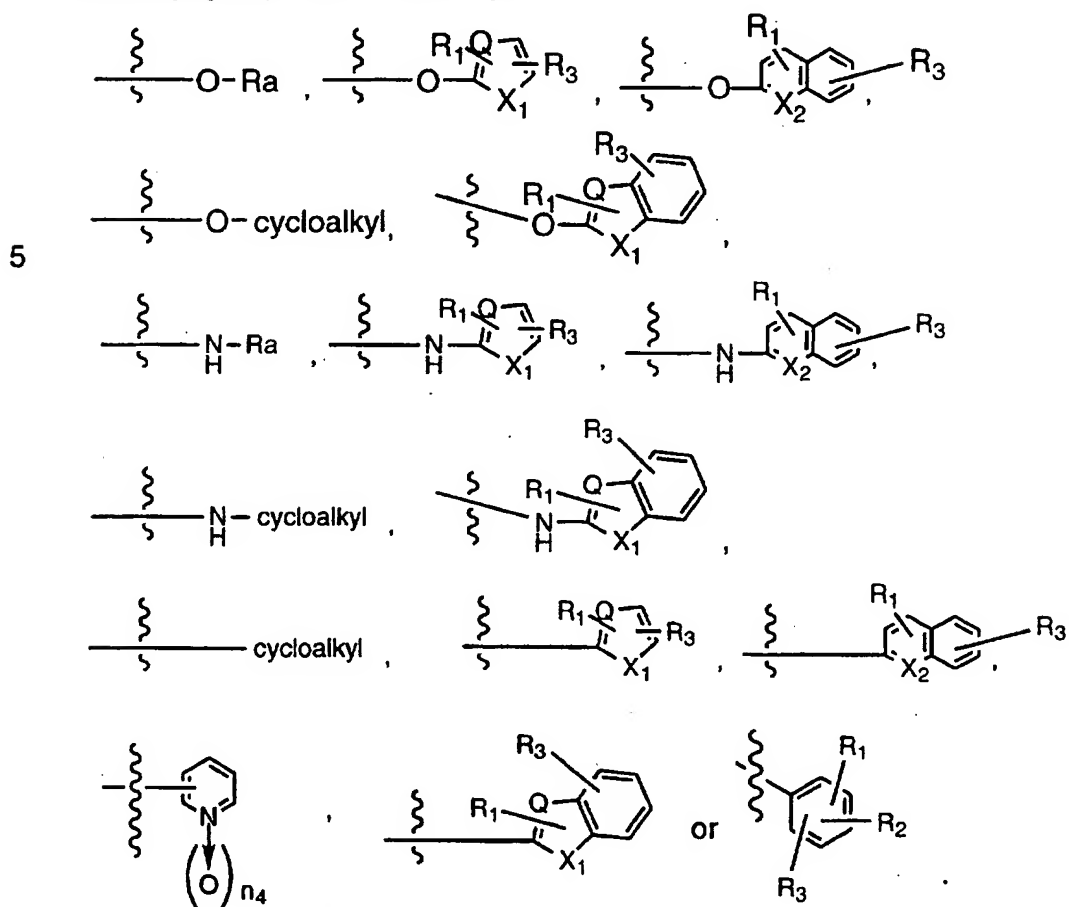
9. A compound according to claim 6 wherein R₆ is



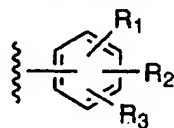
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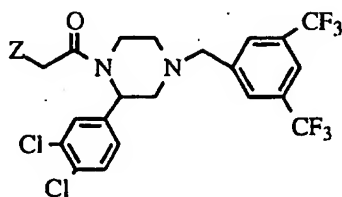
or R_E , R_f taken together with the carbon to which they are attached form a carbonyl group, n_3 is 1, and R_6 is



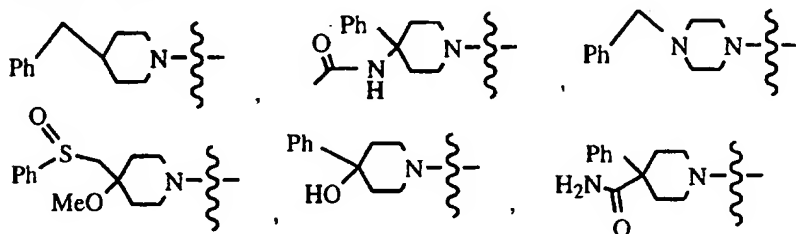
10. A compound according claim 8 or 9 wherein R_6 is



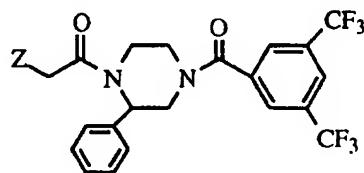
11. A compound according to claim 1 selected from the group consisting of:



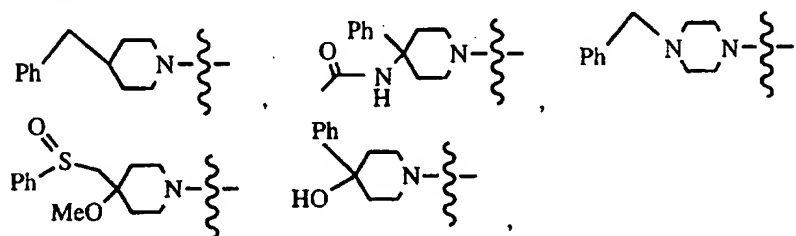
Where Z is



or a compound selected from the group consisting of

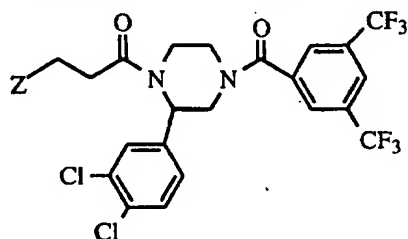


Where Z is

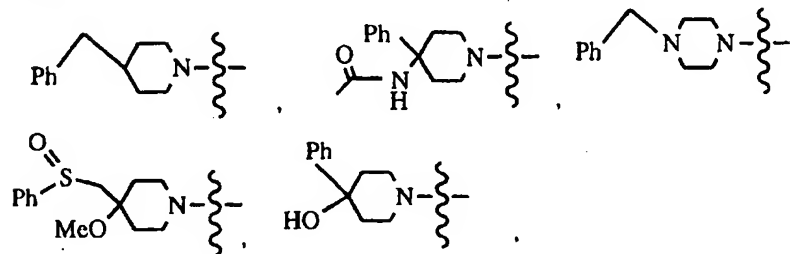


5

or a compound selected from the group consisting of

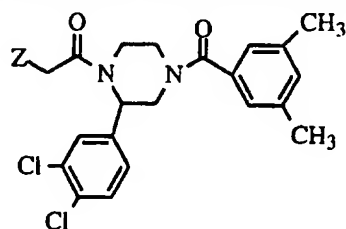


Where Z is

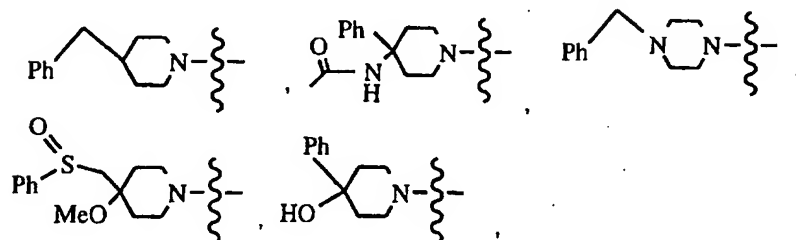


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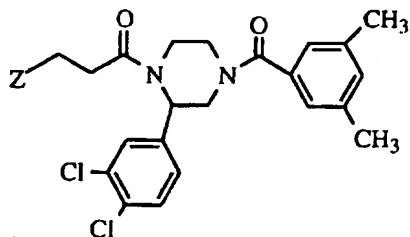
or a compound selected from the group consisting of



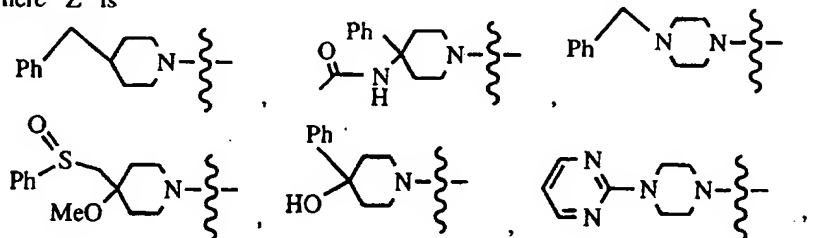
Where Z is



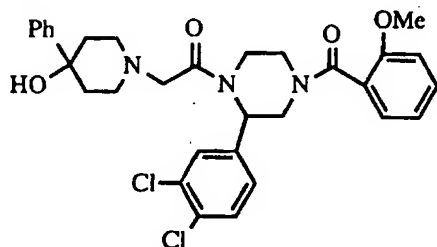
or a compound selected from the group consisting of



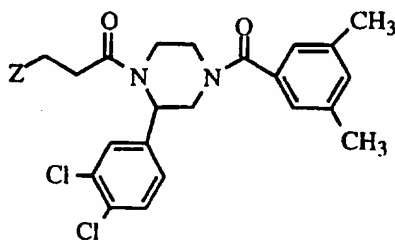
Where Z is



5 or

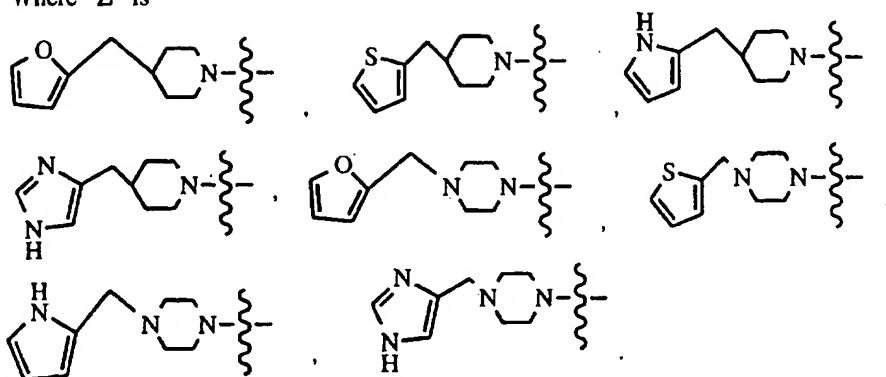


or a compound selected from the group consisting of

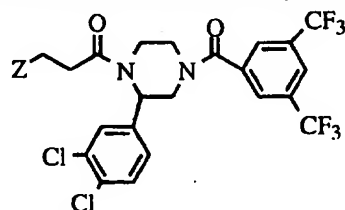


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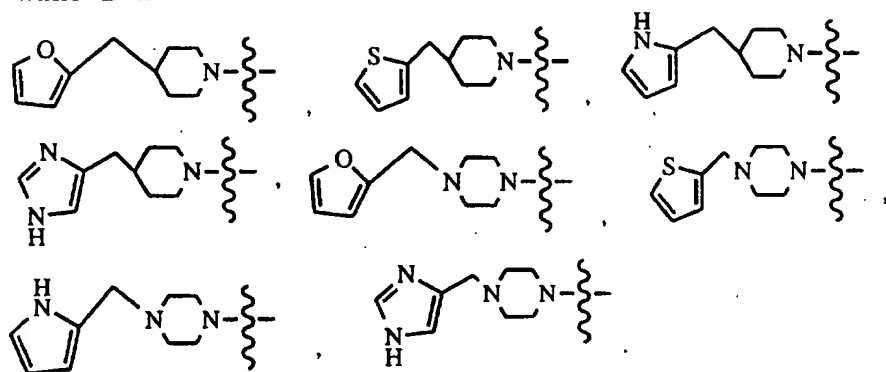
Where Z is



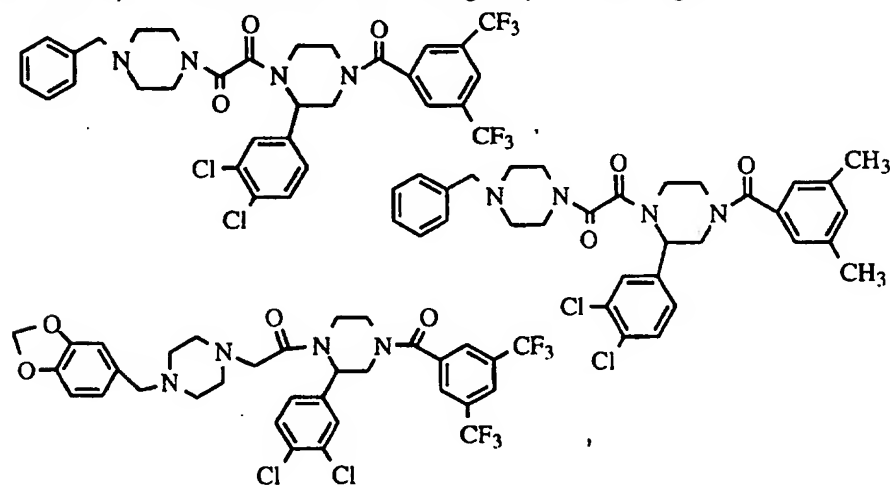
or a compound selected from the group consisting of

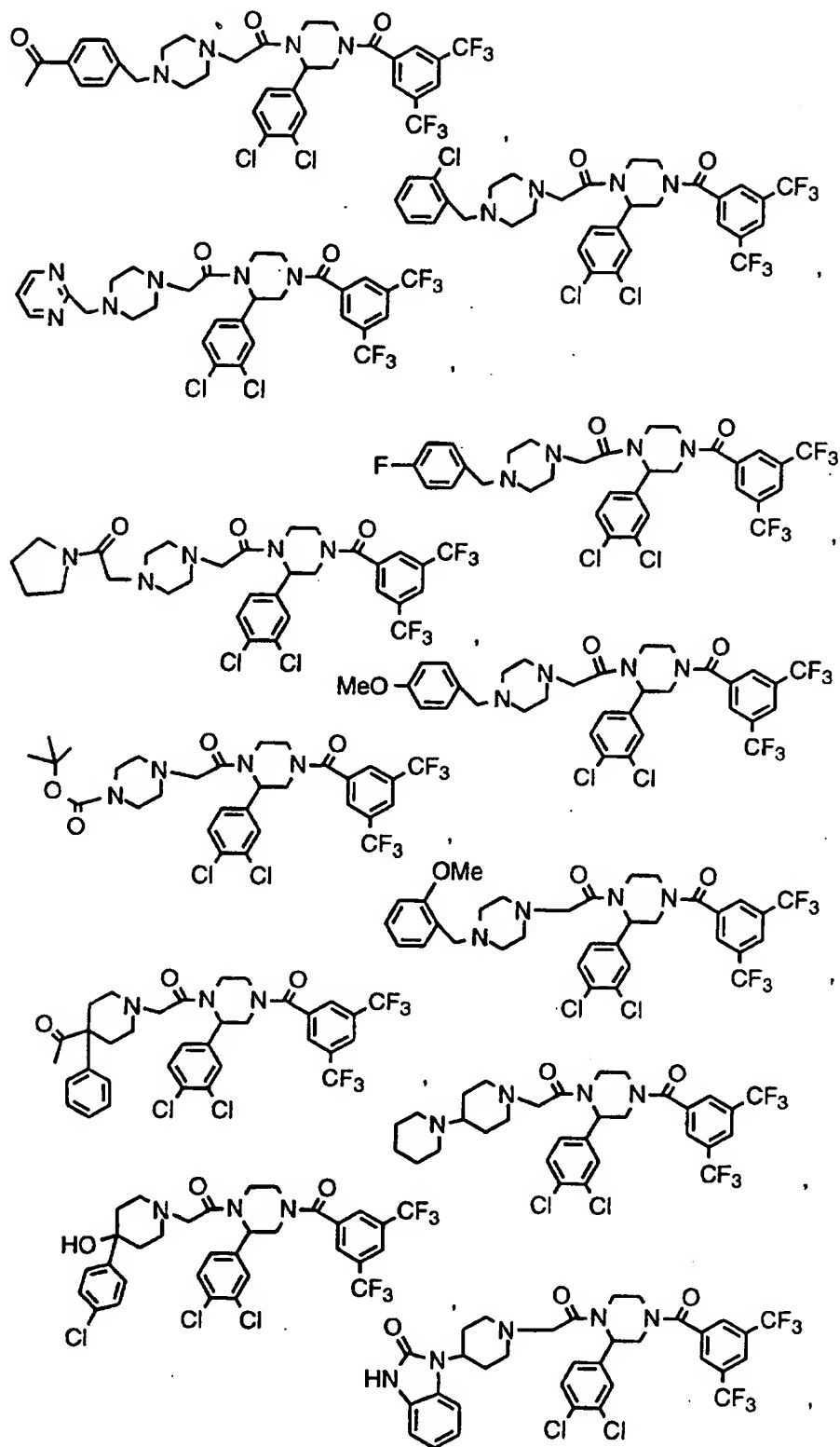


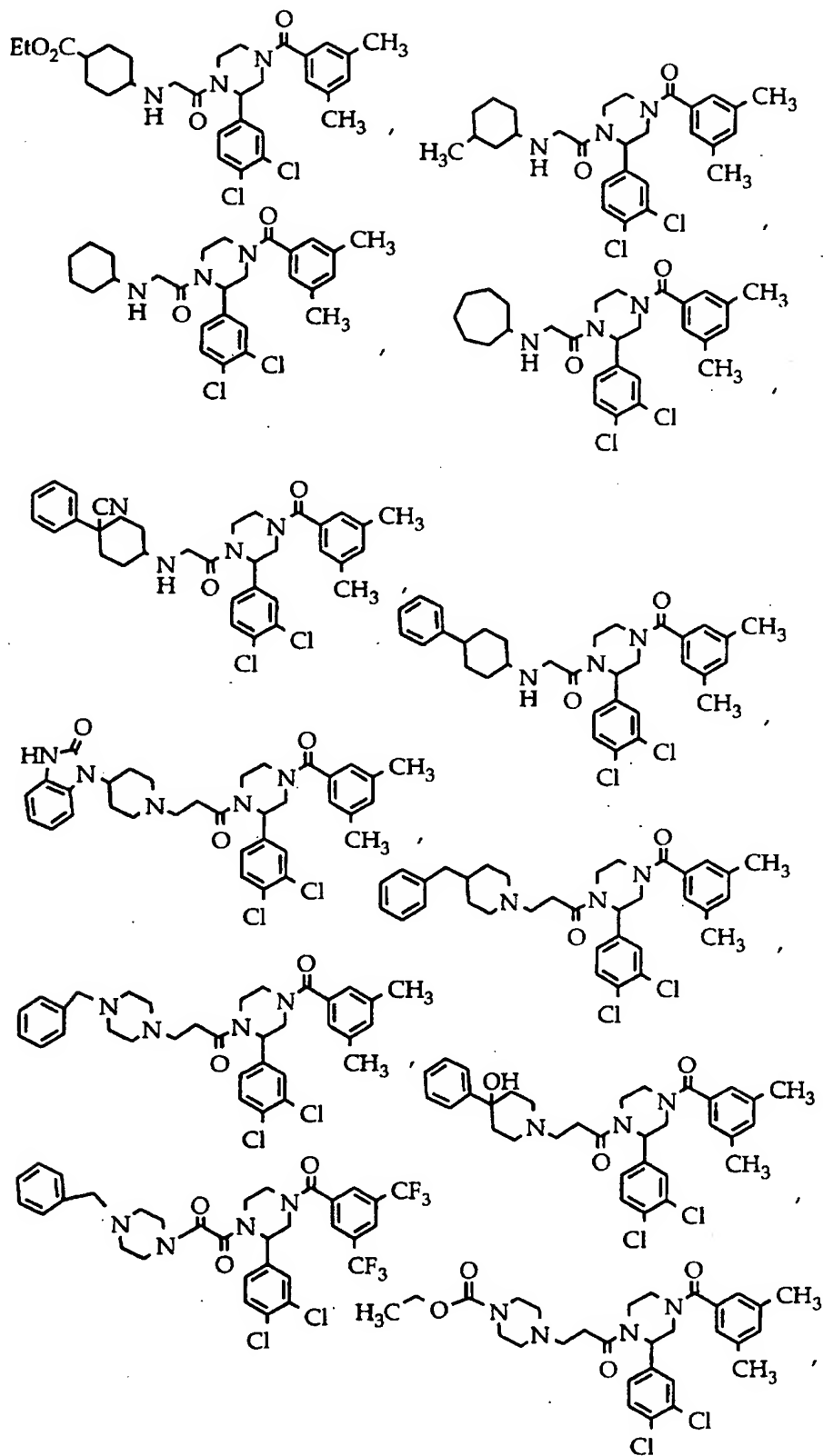
Where Z is

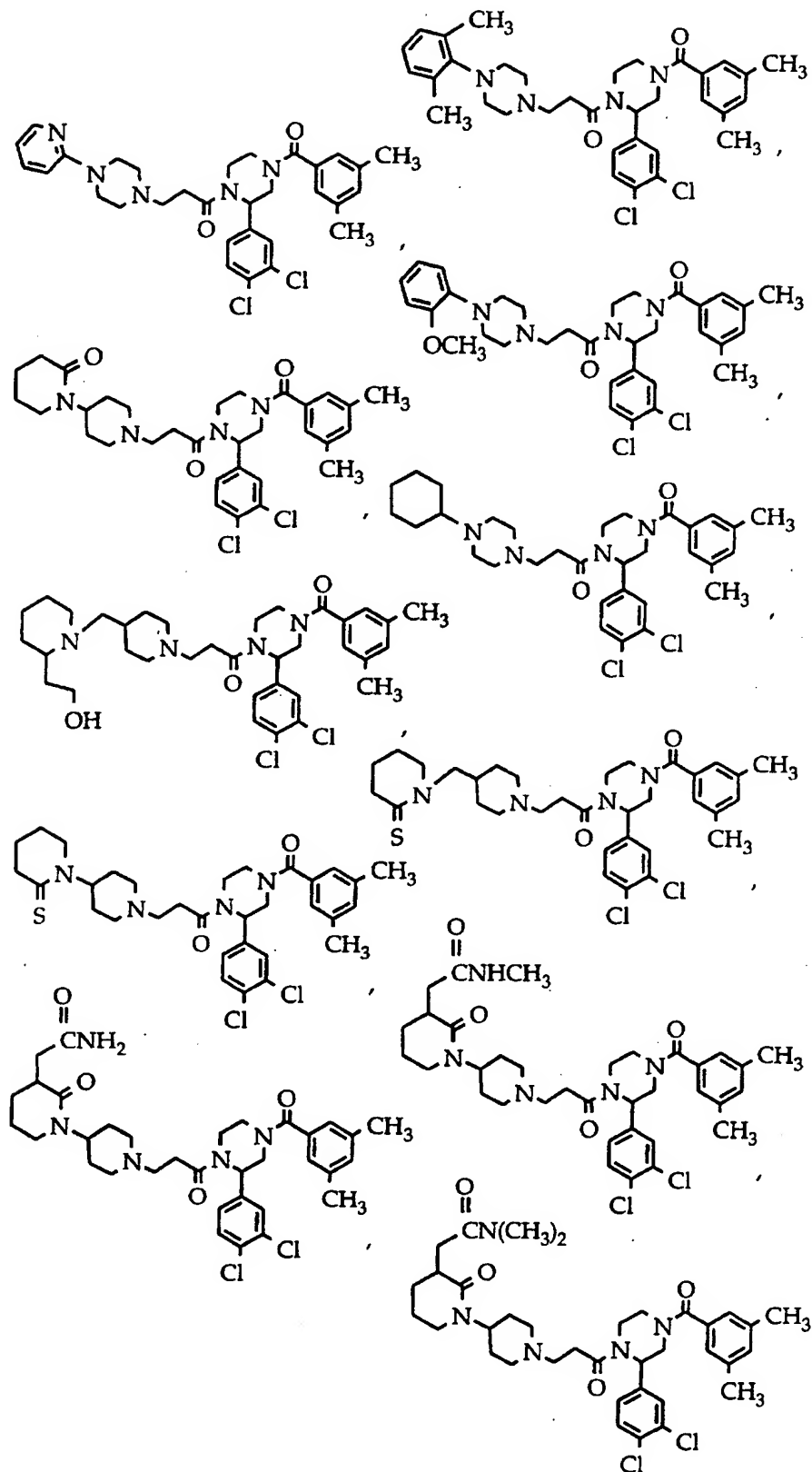


or a compound selected from the group consisting of









or a pharmaceutically acceptable salt thereof.

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12. A pharmaceutical composition comprising a neurokinin antagonistic effective amount of a compound according to any of claims 1 to 11 and a pharmaceutically acceptable carrier material.
- 5 13. The use of a compound of any of claims 1 to 11 for preparing a medicament useful for inducing neurokinin antagonism.
- 10 14. The use of a compound of any of claims 1 to 11 for preparing a medicament useful for treating chronic airway diseases such as asthma, bronchospasm and allergies; inflammatory diseases such as inflammatory bowel disease, psoriasis, fibrositis, osteoarthritis, and rheumatoid arthritis; migraine; central nervous system disorders such as depression, psychosis, dementia, and Alzheimer's disease; Down's syndrome; neuropathy; multiple sclerosis; ophthalmic disorders; conjunctivitis; auto
- 15 immune disorders; graft rejection; systemic lupus erythematosus; GI disorders such as Crohn's disease and ulcerative colitis; disorders of bladder function; circulatory disorders such as angina; Raynaud's disease; emesis, coughing and pain, which comprises administering a therapeutically effective amount of a compound according to claim 1.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 97/14709

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C07D241/04 C07D401/06 C07D405/12 C07D401/14 C07D403/12
A61K31/495

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C07D A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 655 442 A (FUJISAWA PHARMACEUTICAL CO., LTD.) 31 May 1995 see claims ---	1-14
A	WO 94 13646 A (MERCK & CO. INC.) 23 June 1994 see claims ---	1-14
A	US 3 935 214 A (H. ZELLNER) 27 January 1976 see claims ---	1-14
P,X	WO 96 34864 A (SCHERING CORPORATION) 7 November 1996 see the whole document ---	1-14
	-/--	

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

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"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

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"&" document member of the same patent family

Date of the actual completion of the international search

2 December 1997

Date of mailing of the international search report

15/12/1997

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Chouly, J

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 97/14709

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,X	WO 97 08166 A (SCHERING CORPORATION) 6 March 1997 see the whole document -----	1-14

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 97/14709

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